

GenCore version 4.5  
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OM nucleic - nucleic search, using sw model

Run on: July 31, 2002, 18:59:48 ; Search time 2686.26 Seconds  
(Without alignments)  
185.904 Million cell updates/sec

Title: US-09-824-567-4  
Perfect score: 37  
Sequence: 1 ggcgcggatccattttccttagcatacgaagatcc 37

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: EST.\*

1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hic:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
12: gb\_gss:\*  
13: em\_gss\_hum:\*  
14: em\_gss\_inv:\*  
15: em\_gss\_pln:\*  
16: em\_gss\_vrt:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Result No.	Score	Match	Length	DB	ID	Description
1	23.2	62.7	697	12	AZ433593	AZ433593 IM0219H04
2	22.6	61.1	707	12	BH432458	BH432458 BOHAL85TR
3	22	59.5	409	10	B1476980	B1476980 daa87c05
4	22	59.5	419	10	T06394	T06394 EST04283 Fe
5	22	59.5	460	10	BG892973	BG892973 daa92b07
6	22	59.5	480	10	BJ059386	BJ059386 BJ059386
7	22	59.5	573	10	BJ071059	BJ071059 BJ071059
8	22	59.5	575	10	BJ069469	BJ069469 BJ069469
9	22	59.5	616	10	BJ057427	BJ057427 BJ057427
10	22	59.5	618	10	BJ096346	BJ096346 BJ096346
11	21.6	58.4	509	12	TA130r060	TA130r060
12	21.2	57.3	708	12	AQ917850	AQ917850 RPCI-23-2
13	21.2	57.3	741	10	BG432622	BG432622 602500787
14	21	56.8	236	9	AV336182	AV336182 AV336182
15	21	56.8	440	12	AQ517828	AQ517828 HS 5133_B
16	21	56.8	466	12	AQ267958	AQ267958 RPCI11-72
17	21	56.8	556	12	BH063042	BH063042 RPCI-24-3

## SUMMARIES

## ALIGNMENTS

RESULT 1  
AZ433593  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

AZ433593  
LM0219H04R Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
clone UUGC1M0219H04 R, DNA sequence.  
697 bp DNA linear GSS 03-OCT-2000  
AZ433593  
GSS.  
GI:10557606  
house mouse.  
Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.  
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,  
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,  
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.  
and Wright, D., Weiss, R.  
Mouse whole genome scaffolding with paired end reads from 10kb  
plasmid inserts  
Unpublished (2000)  
Contact: Robert B. Weiss  
University of Utah Genome Center  
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunn@genetics.utah.edu  
Insert Length: 10000 Std Error: 0.00  
Plate: 0219 row: H column: 04  
Seq primer: CACACAGGAACAGCTATGACC  
Class: plasmid ends  
High quality sequence stop: 697.  
Location/Qualifiers  
1. .697  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UUGC1M0219H04"  
/clone\_ltb="Mouse 10kb plasmid UUGC1M library"

FEATURES  
source

/sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
 /note="Vector: PWD42nv: Purified genomic DNA from M.  
 musculus C57BL/6J (male) was obtained from the Jackson  
 Laboratory Mouse DNA Resource  
 (http://www.jax.org/resources/documents/dnares/). The DNA  
 was hydrodynamically sheared by repeated passage through a  
 0.005 inch orifice at constant velocity. The sheared DNA  
 was blunt end-repaired with T4 DNA polymerase and T4  
 polynucleotide kinase. Adaptor oligonucleotides were  
 ligated to the blunt ends in high molar excess. The  
 adaptor DNA was purified and size-selected for a 9.5 to  
 10.5 kb range using preparative agarose gel  
 electrophoresis. Vector DNA was prepared from a derivative  
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number  
 inducible derivative of plasmid R1. The vector was ligated  
 with adaptors complementary to the insert adaptors and  
 purified. The sheared, adaptor mouse DNA was annealed to  
 adaptor vector DNA, and transformed into  
 chemically-competent E. coli XL10-Gold (Stratagene) cells  
 and selected for ampicillin resistance."

BASE COUNT 192 a 189 c 104 g 212 t  
 ORIGIN

Query Match 62.7%; Score 23.2; DB 12; Length 697;  
 Best Local Similarity 77.8%; Pred. No. 30;  
 Matches 28; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 1 gcgcggatccatttccttagcataacgaagtc 36  
 Db 16 GTGCCGAATCCTACTTCCTTCCTTGCCTATGGAATC 51

RESULT 2  
 BH432458 707 bp DNA linear GSS 12-DEC-2001  
 LOCUS BOHAL85TR BOHA Brassica oleracea genomic clone BOHAL85, DNA  
 DEFINITION  
 ACCESSION BH432458  
 VERSION BH432458  
 KEYWORDS GSS.  
 SOURCE Brassica oleracea.  
 ORGANISM Brassica oleracea

REFERENCE 1. (bases 1 to 707)  
 AUTHORS Town, C.D., Van Aken, S., Utterback, T. and Fraser, C.M.  
 TITLE Whole genome shotgun sequencing of Brassica oleracea  
 JOURNAL Unpublished (2001)  
 COMMENT Other GSSs: BOHAL85TF  
 Contact: Chris Town  
 TIGR

7712 Medical Center Drive, Rockville, MD 20850, USA.  
 Tel: 301-838-3523  
 Fax: 301-838-0208  
 Email: cdtown@tigr.org  
 DNA is from a doubled haploid provided by Tom Osborn.  
 Seq primer: TR  
 Class: sheared ends.

FEATURES  
 source

1. .707  
 /organism="Brassica oleracea"  
 /strain="T01000DH3"  
 /db\_xref="taxon:3712"  
 /clone="BOHAL85"  
 /note="Vector: pBstXI; Site 1: BstXI; 2-3 kb sheared  
 genomic DNA inserted into pBstXI using BstXI linkers"

BASE COUNT 245 a 139 c 131 g 192 t  
 ORIGIN

Query Match 61.1%; Score 22.6; DB 12; Length 707;  
 Best Local Similarity 86.2%; Pred. No. 54;  
 Matches 25; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 8 atccatttccttagcataacgaagtc 36  
 Db 215 ATACATTTTCCTAGCAAAACGTAAGTC 243

RESULT 3  
 BI476980/4 409 bp mRNA linear EST 27-AUG-2001  
 LOCUS daa87c05.y4 Wellcome CRC PRN3 St13 17 egg animal cap Xenopus laevis  
 DEFINITION  
 ACCSSION BI476980  
 VERSION BI476980  
 KEYWORDS EST.  
 SOURCE African clawed frog.  
 ORGANISM Xenopus laevis

REFERENCE 1. (bases 1 to 409)  
 AUTHORS Clifton, S., Johnson, S.L., Blumberg, B., Song, J., Hillier, L., Pape, D.,  
 B., Gibbons, M., Harvey, N., Ritter, E., Jackson, Y., McCann, R.,  
 Waterston, R. and Wilson, R.  
 TITLE WashU xenopus EST project, 1999  
 JOURNAL Unpublished (1999)  
 COMMENT Other ESTs: daa87c05.x3  
 Contact: Sandy Clifton, Ph.D.  
 WashU xenopus EST project, 1999  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@watson.wustl.edu

Library constructed by N. Garrett, E. ellefroid, and A.M. Zorn  
 (Wellcome/CRC Institute). DNA Sequencing by: Washington University  
 Genome Sequencing Center  
 Clone distribution: Xenopus clones from this library are available  
 through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov  
 High quality sequence stop: 383.  
 Location/Qualifiers  
 1. .409  
 /organism="Xenopus laevis"  
 /db\_xref="taxon:8355"  
 /clone="IMAGE:4084232"  
 /clone\_lib="Wellcome CRC PRN3 St13 17 egg animal cap"  
 /tissue\_type="egg, subtracted by stage 13-17 animal cap"  
 /lab\_host="DHI09 (phage-resistant)"  
 /note="Vector: pBSRN3; Site 1: NotI; Site 2: EcoRI; cDNAs  
 were oligo-dT primed and directionally cloned. Staging  
 according to Nieuwkoop and Faber. Library is subtracted  
 and was constructed by N. Garrett, E. ellefroid, and A.M.  
 Zorn, (Wellcome/CRC Institute)."

BASE COUNT 94 a 94 c 108 g 113 t  
 ORIGIN

Query Match 59.5%; Score 22; DB 10; Length 409;  
 Best Local Similarity 83.3%; Pred. No. 87;  
 Matches 25; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 4 ccgagatccatttccttagcataacgaa 33  
 Db 278 CGGATCCCATTCCTCCACAGCAGATGGA 249

RESULT 4  
 T06394/c  
 LOCUS T06394

419 bp mRNA linear EST 30-JUN-1993

```

Query Match          59.5%; Score 22; DB 10; Length 480;
Best Local Similarity 83.3%; Pred. No. 89;
Matches 25; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

```

Qy 4 ccggatcccatcttctccttagcataacggaa 33  
|||||  
Db 77 CCGGATCCCATCTCTCCACAGCAGATGGAA 106  
|||||

## RESULTS

R28911	/
BJ071059	
LOCUS	
DEFINITION	Bj071059 NIBB Mochii normalized Xenopus tailbud library Xenopus laevis cDNA clone XL092n13 5' mRNA sequence.
ACCESSION	X071059
VERSION	BJ071059
DATE	EST 11-DEC-2001

KEYWORDS  
E5071033.1 G1:1/501248  
F57

SOURCE	African clawed frog.
ORGANISM	Xenopus laevis
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
	Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
REFERENCE	Xenopodinae Xenopus
	1 (bases 1 to 575)
AUTHORS	Kitayama, A., Terasaka, C., Mochii, M., Ueno, N., Shin-i, T. and Kohara

1. Expressed genes in *X. laevis* embryo

Unpublished (2001)  
Contact: Tadasu Shin-i  
Center For Genetic Resource Information  
National Institute of Genetics  
1111 Yats, Mishima, Shizuoka 411-8540, Japan  
Tel: 81-559-81-6856  
Fax: 81-559-81-6855

## FEATURES

FEATURES	Location/Qualifiers
source	1. .573

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/organism="Xenopus laevis"  
/db_xref="taxon:8355"  
/clone="XL092n13"  
/clone.lib="NIBB Mochii normalized Xenopus tailbud  
library"
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BASE COUNT	136 a	134 c	123 g	180 t
ORIGIN				

Query Match 59.5%; Score 22; DB 10; Length 573;  
Best Local Similarity 83.3%; Pred. NO. 92;  
Matches 25; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

4 cggatcccat<sup>1</sup>ttccttagcataacqqa<sup>2</sup> 33

[illegible]

ACCESSION BJ069469

VERSION  
 WORDS  
 SYNONYMS  
 SOURCE  
 ORIGIN  
 ORGANISM  
 EST.  
 BT069469.1 GI:17497829  
 African clawed frog.  
 Xenopus laevis  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;  
 Xenopodinae; Xenopus.  
 1 (bases 1 to 575)  
 Kitayama, A., Terasaka, C., Mochii, M., Ueno, N., Shin-i, T. and Kobara  
 Y.

TITLE Expressed genes in *X. laevis* embryo

Unpublished (2001)  
Contact: Tadasu Shin-i  
Center For Genetic Resource Information  
National Institute of Genetics

1111 Yata, Mishima, Shizuoka 411-8540, Japan  
Tel: 81-559-81-6856  
Fax: 81-559-81-6855  
Email: tshiniegenes.nig.ac.jp.

FEATURES	SOURCE
1. <i>General</i>	
2. <i>Specific</i>	
3. <i>Other</i>	
4. <i>Other</i>	
5. <i>Other</i>	
6. <i>Other</i>	
7. <i>Other</i>	
8. <i>Other</i>	
9. <i>Other</i>	
10. <i>Other</i>	
11. <i>Other</i>	
12. <i>Other</i>	
13. <i>Other</i>	
14. <i>Other</i>	
15. <i>Other</i>	
16. <i>Other</i>	
17. <i>Other</i>	
18. <i>Other</i>	
19. <i>Other</i>	
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21. <i>Other</i>	
22. <i>Other</i>	
23. <i>Other</i>	
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26. <i>Other</i>	
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32. <i>Other</i>	
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59. <i>Other</i>	
60. <i>Other</i>	
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63. <i>Other</i>	
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65. <i>Other</i>	
66. <i>Other</i>	
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70. <i>Other</i>	
71. <i>Other</i>	
72. <i>Other</i>	
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76. <i>Other</i>	
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91. <i>Other</i>	
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96. <i>Other</i>	
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100. <i>Other</i>	

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i: 3373
/organism="Xenopus laevis"
/db_xref="taxon:8355"
/clone="XI053a06"
/clone_lib="NIBB Mochii normalized Xenopus tailbud
library"
/tissue_type="whole embryo"

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BASE COUNT	/dev_stage="stage 25"
162 a	132 c 136 g
ORIGIN	143 t 2 others

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Query Match          59.5%; Score 22; DB 10; Length 575;
Best Local Similarity 83.3%; Pred. No. 92;
Matches 25; Count 1;

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Qy 4 ccggatcccattttccttagcataacgaa 33  
|||||  
Db 223 CCGGATCCCATTCTCCACACGAGAATGGAA 252

PRESENT

	BJ057427	LOCUS	616 bp mRNA linear EST 10-DEC-2001
	Bj057427	NIBB Mochii normalized Xenopus tailbud library Xenopus laevis cDNA clone XL10441 5' mRNA sequence.	
RESULTS	Bj057427	DEFINITION	
	Bj057427	ACCESSION	

ACCESSION B057427

BJ057427.1 GI:17470221  
EST.  
SOURCE  
ORGANISM  
Xenopus laevis  
African clawed frog.  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;  
Xenopodinae; Xenopus.  
1 (bases 1 to 616)  
Kitayama,A., Terasaka,C., Mochii,M., Ueno,N., Shin-I,T. and Kohara  
Y.

**TITLE** Expressed genes in *X. laevis* embryo  
**JOURNAL** Unpublished (2001)  
**COMMENT** Contact: Tadasu Shin-i  
Center For Genetic Resource Information  
National Institute of Genetics  
1111 Yatai, Mishima, Shizuoka 411-8540, Japan  
Tel: 81-559-81-6856  
Fax: 81-559-81-6855

**REFERENCES**

```

Location/Qualifiers
1. .616
/organism="Xenopus laevis"
/db_xref="taxon:8355"
/clone="Xrl04all"
/clone_lib="NIBB Mochii normalized Xenopus tailbud
library"
/tissue_type="whole embryo"

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BASE COUNT	/dev_stage="stage 25"	115 t	2 others
188 a	150 c	161 g	
188 a	150 c	161 g	

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Best Local Similarity	83.3%;	Pred. No. 93;		
Matches	25			

4 ccggtatcccatctttccttagcataacgga 33  
|||||  
399 CCGGATCCCATCTTCTCCACAGCAGAATGGA 428

conducted at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of *Trypanosoma brucei* (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (4 kb). The *v* + i method used for the library construction is





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OM nucleic - nucleic search, using sw model

Run on: July 31, 2002, 19:32:59 ; Search time 84.08 Seconds  
(without alignments)  
108.093 Million cell updates/sec

Title: US-09-824-567-4

Perfect score: 37

Sequence: 1 ggcgcggatccatttccttagcatacgaagtc 37

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 383533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued\_Patents.NA.\*  
1: /cgn2\_6/ptodata/2/ina/5A\_COMB.seq.\*  
2: /cgn2\_6/ptodata/2/ina/5B\_COMB.seq.\*  
3: /cgn2\_6/ptodata/2/ina/6A\_COMB.seq.\*  
4: /cgn2\_6/ptodata/2/ina/6B\_COMB.seq.\*  
5: /cgn2\_6/ptodata/2/ina/PTCUS\_COMB.seq.\*  
6: /cgn2\_6/ptodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	19.8	53.5	497	3	US-09-248-528-4
2	19.8	53.5	497	4	US-09-549-108-4
3	19.8	53.5	497	4	US-09-549-111-4
4	19.8	53.5	497	4	US-09-549-106-4
5	19.8	53.5	497	4	US-09-550-394-4
6	18.2	49.2	10803	3	US-09-080-044-1
7	17.8	48.1	30	4	US-09-522-666-13
8	17.8	48.1	2859	3	US-09-170-354-7
9	17.8	48.1	2859	3	US-09-522-666-1
10	17.6	47.6	3092	4	US-09-522-666-1
11	17.4	47.0	6464	1	US-08-321-478-2
12	17.4	47.0	6464	1	US-08-321-478-4
13	17.4	47.0	6464	1	US-08-321-478-6
14	17.4	47.0	10754	2	US-08-966-958-1
15	17.4	47.0	10754	2	US-09-215-817-1
16	17.4	47.0	10754	2	US-09-342-353-1
17	17.4	47.0	11236	1	US-07-853-913-1
18	17.2	46.5	593	4	US-09-328-111-724
19	17.2	46.5	607	3	US-08-894-483-6
20	17.2	46.5	4895	4	US-09-426-568A-3
21	17	45.9	573	4	US-09-385-982-451
22	17	45.9	830	4	US-08-998-416-298
23	17	45.9	1460	1	US-08-133-038A-1
24	17	45.9	1460	1	US-08-161-988A-1
25	17	45.9	2149	1	US-08-784-651-3
26	17	45.9	8752	4	US-08-976-259-3
27	17	45.9	72928	3	US-09-009-913-1

28 16.8 45.4 566 1 US-08-663-023-16  
29 16.8 45.4 2362 1 US-08-265-087-1  
30 16.8 45.4 2362 1 US-08-621-493-1  
31 16.8 45.4 2362 2 US-08-965-688-1  
32 16.8 45.4 2362 4 US-09-260-173-1  
33 16.8 45.4 2458 3 US-08-611-587-6  
34 16.8 45.4 2994 1 US-08-204-329-2  
35 16.8 45.4 2994 2 US-08-482-627-4  
36 16.8 45.4 2994 3 US-08-801-092-3  
37 16.8 45.4 2994 5 PCT-US94-10357-1  
38 16.8 45.4 2995 2 US-08-959-638-7  
39 16.8 45.4 2995 4 US-08-328-673A-7  
40 16.8 45.4 3153 4 US-09-175-928-9  
41 16.8 45.4 3232 1 US-08-038-760-1  
42 16.8 45.4 3232 1 US-08-038-760-2  
43 16.8 45.4 3232 2 US-08-470-091-1  
44 16.8 45.4 3232 2 US-08-470-091-2  
45 16.8 45.4 3300 3 US-08-913-842-4

## ALIGNMENTS

RESULT 1  
US-09-248-528-4  
; Sequence 4, Application US/09248528C  
; Patent No. 6153415  
; GENERAL INFORMATION:  
; APPLICANT: Oriel, Patrick J  
; APPLICANT: Padmakumar, Rugmini  
; APPLICANT: Kim, Sang H  
; TITLE OF INVENTION: Method for Producing Amide Compounds Using a Nitrile  
; TITLE OF INVENTION: Hydrtase from a Thermophilic Bacillus  
; FILE REFERENCE: MSU 4.1-401  
; CURRENT APPLICATION NUMBER: US/09/248,528C  
; CURRENT FILING DATE: 1999-02-10  
; EARLIER APPLICATION NUMBER: 60/083,485  
; EARLIER FILING DATE: 1998-04-29  
; NUMBER OF SEQ ID NOS: 18  
; SOFTWARE: Patentin Ver. 2.0  
; SEQ ID NO 4  
; LENGTH: 497  
; TYPE: DNA  
; ORGANISM: Bacillus smithii  
; FEATURE:  
; NAME/KEY: rRNA  
; LOCATION: (1)..(497)  
; OTHER INFORMATION: nucleotides 17 - 513 of rRNA sequence  
; OTHER INFORMATION: X60643/Genbank  
; PUBLICATION INFORMATION:  
; DATABASE ACCESSION NUMBER: X60643/Genbank  
; DATABASE ENTRY DATE: 1997-04-03  
US-09-248-528-4

Query Match 53.5%; Score 19.8; DB 3; Length 497;  
Best Local Similarity 77.4%; Pred. No. 5.1;  
Matches 24; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 4 ccggatccatttccttagcatacgaag 34  
||||| ||||||| |||||  
Db 152 ccggataatactcttcgcaggaag 182

RESULT 2  
US-09-549-108-4  
; Sequence 4, Application US/09549108  
; Patent No. 6214603  
; GENERAL INFORMATION:  
; APPLICANT: Oriel, Patrick J  
; APPLICANT: Padmakumar, Rugmini  
; APPLICANT: Kim, Sang H  
; TITLE OF INVENTION: Method for Producing Amide Compounds Using a Nitrile



; TITLE OF INVENTION: Hydratase from a Thermophilic Bacillus  
; FILE REFERENCE: MSU 4.1-486  
; CURRENT APPLICATION NUMBER: US/09/549,108  
; CURRENT FILING DATE: 2000-04-13  
; PRIOR APPLICATION NUMBER: 60/083,485  
; PRIOR FILING DATE: 1998-04-29  
; PRIOR APPLICATION NUMBER: 09/248,528  
; PRIOR FILING DATE: 1999-02-10  
; NUMBER OF SEQ ID NOS: 18  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 4  
; LENGTH: 497  
; TYPE: DNA  
; ORGANISM: Bacillus smithii  
; FEATURE:  
; NAME/KEY: rRNA  
; LOCATION: (1)..(497)  
; OTHER INFORMATION: nucleotides 17 - 513 of rRNA sequence  
; OTHER INFORMATION: X60643/Genbank  
; PUBLICATION INFORMATION:  
; DATABASE ACCESSION NUMBER: X60643/Genbank  
; DATABASE ENTRY DATE: 1997-04-03  
US-09-549-108-4

Query Match 53.5%; Score 19.8; DB 4; Length 497;  
Best Local Similarity 77.4%; Pred. No. 5.1;  
Matches 24; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 4 ccggaatccattctcttagcatacgaag 34  
||||| || ||||| ||||| |||||  
DB 152 ccggaataatctcttcgcgaaggaag 182

## RESULT 3

US-09-549-111-4  
; Sequence 4, Application US/09549111  
; Patent No. 6228633

; GENERAL INFORMATION:  
; APPLICANT: Oriel, Patrick J  
; APPLICANT: Padmakumar, Rugmini  
; APPLICANT: Kim, Sang H  
; TITLE OF INVENTION: Method for Producing Amide Compounds Using a Nitrile  
; TITLE OF INVENTION: Hydratase from a Thermophilic Bacillus  
; FILE REFERENCE: MSU 4.1-489  
; CURRENT APPLICATION NUMBER: US/09/549,111  
; PRIOR APPLICATION NUMBER: 60/083,485  
; PRIOR FILING DATE: 1998-04-29  
; PRIOR APPLICATION NUMBER: 09/248,528  
; PRIOR FILING DATE: 1999-02-10  
; NUMBER OF SEQ ID NOS: 18  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 4  
; LENGTH: 497  
; TYPE: DNA  
; ORGANISM: Bacillus smithii  
; FEATURE:  
; NAME/KEY: rRNA  
; LOCATION: (1)..(497)  
; OTHER INFORMATION: nucleotides 17 - 513 of rRNA sequence  
; OTHER INFORMATION: X60643/Genbank  
; PUBLICATION INFORMATION:  
; DATABASE ACCESSION NUMBER: X60643/Genbank  
; DATABASE ENTRY DATE: 1997-04-03  
US-09-549-111-4

Query Match 53.5%; Score 19.8; DB 4; Length 497;  
Best Local Similarity 77.4%; Pred. No. 5.1;  
Matches 24; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 4 ccggaatccattctcttagcatacgaag 34

Db 152 ccggaataatctcttcgcgaaggaag 182  
||||| || ||||| ||||| |||||

## RESULT 4

US-09-549-106-4  
; Sequence 4, Application US/09549106  
; Patent No. 6242242

; GENERAL INFORMATION:  
; APPLICANT: Oriel, Patrick J  
; APPLICANT: Padmakumar, Rugmini  
; APPLICANT: Kim, Sang H  
; TITLE OF INVENTION: Method for Producing Amide Compounds Using a Nitrile  
; TITLE OF INVENTION: Hydratase from a Thermophilic Bacillus  
; FILE REFERENCE: MSU 4.1-487  
; CURRENT APPLICATION NUMBER: US/09/549,106  
; CURRENT FILING DATE: 2000-04-13  
; PRIOR APPLICATION NUMBER: 60/083,485  
; PRIOR FILING DATE: 1998-04-29  
; PRIOR APPLICATION NUMBER: 09/248,528  
; PRIOR FILING DATE: 1999-02-10  
; NUMBER OF SEQ ID NOS: 18  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 4  
; LENGTH: 497  
; TYPE: DNA  
; ORGANISM: Bacillus smithii  
; FEATURE:  
; NAME/KEY: rRNA  
; LOCATION: (1)..(497)  
; OTHER INFORMATION: nucleotides 17 - 513 of rRNA sequence  
; OTHER INFORMATION: X60643/Genbank  
; PUBLICATION INFORMATION:  
; DATABASE ACCESSION NUMBER: X60643/Genbank  
; DATABASE ENTRY DATE: 1997-04-03  
US-09-549-106-4

Query Match 53.5%; Score 19.8; DB 4; Length 497;  
Best Local Similarity 77.4%; Pred. No. 5.1;  
Matches 24; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 4 ccggaatccattctcttagcatacgaag 34  
||||| || ||||| ||||| |||||  
DB 152 ccggaataatctcttcgcgaaggaag 182

## RESULT 5

US-09-550-394-4  
; Sequence 4, Application US/09550394  
; Patent No. 6267829

; GENERAL INFORMATION:  
; APPLICANT: Oriel, Patrick J  
; APPLICANT: Padmakumar, Rugmini  
; APPLICANT: Kim, Sang H  
; TITLE OF INVENTION: Method for Producing Amide Compounds Using a Nitrile  
; TITLE OF INVENTION: Hydratase from a Thermophilic Bacillus  
; FILE REFERENCE: MSU 4.1-488  
; CURRENT APPLICATION NUMBER: US/09/550,394  
; CURRENT FILING DATE: 2000-04-14  
; PRIOR APPLICATION NUMBER: 60/083,485  
; PRIOR FILING DATE: 1998-04-29  
; PRIOR APPLICATION NUMBER: 09/248,528  
; PRIOR FILING DATE: 1999-02-10  
; NUMBER OF SEQ ID NOS: 18  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 4  
; LENGTH: 497  
; TYPE: DNA  
; ORGANISM: Bacillus smithii  
; FEATURE:  
; NAME/KEY: rRNA  
; LOCATION: (1)..(497)

; OTHER INFORMATION: nucleotides 17 - 513 of rRNA sequence  
; PUBLICATION INFORMATION: X60643/Genbank  
; DATABASE ACCESSION NUMBER: X60643/Genbank  
; DATABASE ENTRY DATE: 1997-04-03  
US-09-550-394-4

Query Match 53.5%; Score 19.8; DB 4; Length 497;  
Best Local Similarity 77.4%; Pred. No. 5.1;  
Matches 24; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 4 ccgcatccatttcttagcataacggaag 34  
||||| ||||||| ||||| |||||  
Db 152 ccggaataatcttcttcgaatgaagaa 182

RESULT 6  
US-09-080-044-1  
; Sequence 1, Application US/09080044  
; Patent No. 6074649  
; GENERAL INFORMATION:  
; APPLICANT: AUDONNET, Jean-Christophe F.  
; APPLICANT: BAUDU, Philippe G.  
; APPLICANT: RIVIERE, Michel A.  
; TITLE OF INVENTION: RECOMBINANT VACCINE CONTAINING FELINE HERPES VIRUS TYPE  
; TITLE OF INVENTION: 1, PARTICULARLY FOR TREATING FELINE INFECTIOUS  
; FILE REFERENCE: PERTONITIS  
; CURRENT APPLICATION NUMBER: US/09/080,044  
; CURRENT FILING DATE: 1998-05-15  
; EARLIER APPLICATION NUMBER: PCT/FR96/01830  
; EARLIER FILING DATE: 1996-11-19  
; EARLIER APPLICATION NUMBER: 95/14450  
; EARLIER FILING DATE: 1995-11-30  
; NUMBER OF SEQ ID NOS: 33  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 1  
; LENGTH: 10803  
; TYPE: DNA  
; ORGANISM: Feline herpesvirus 1  
US-09-080-044-1

Query Match 49.2%; Score 18.2; DB 3; Length 10803;  
Best Local Similarity 74.2%; Pred. No. 53;  
Matches 23; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 3 gccgatccatttcttagcataacgga 33  
||||| ||||||| ||||| |||||  
Db 310 gccgatccatttcttagcataacgga 340

RESULT 7  
US-09-522-666-13  
; Sequence 13, Application US/09522666  
; Patent No. 6333167  
; GENERAL INFORMATION:  
; APPLICANT: Shuey, David  
; APPLICANT: Quinet, Blaine  
; TITLE OF INVENTION: Methods and Reagents for Identifying Inhibitors of  
; FILE REFERENCE: 6-00  
; CURRENT APPLICATION NUMBER: US/09/522,666  
; CURRENT FILING DATE: 2000-03-10  
; NUMBER OF SEQ ID NOS: 32  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 13  
; LENGTH: 30  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence:oligonucleotide

US-09-522-666-13

Query Match 48.1%; Score 17.8; DB 4; Length 30;  
Best Local Similarity 75.9%; Pred. No. 21;  
Matches 22; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 1 ggcggatccatttcttagcataac 29  
||||| ||||||| ||||| |||||  
Db 1 gtcggatccatttcttagcataac 29

RESULT 8  
US-08-637-763B-7  
; Sequence 7, Application US/08637763B  
; Patent No. 5849559  
; GENERAL INFORMATION:  
; APPLICANT: VAN DER WOUW, Monique J.A. et al  
; TITLE OF INVENTION: ARABINOXYLAN DEGRADING ENZYME  
; NUMBER OF SEQUENCES: 8  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Morrison & Foerster  
; STREET: 2000 Pennsylvania Avenue, NW  
; CITY: Washington  
; STATE: DC  
; COUNTRY: USA  
; ZIP: 20006-1812  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/637,763B  
; FILING DATE: 25-AUG-1996  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Murashige, Kate H.  
; REGISTRATION NUMBER: 29,959  
; REFERENCE/DOCKET NUMBER: 4615-0066.00  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (202) 887-1500  
; TELEFAX: (202) 887-0763  
; TELEX: 90-4030 MRSNFORSHW  
; INFORMATION FOR SEQ ID NO: 7:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2859 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; HYPOTHETICAL: NO  
; ANTI-SENSE: NO  
; ORIGINAL SOURCE:  
; ORGANISM: Aspergillus niger var. tubigenis  
; STRAIN: DS16813  
; FEATURE:  
; NAME/KEY: CAAT\_signal  
; LOCATION: 651..655  
; FEATURE:  
; NAME/KEY: TATA\_signal  
; LOCATION: 713..720  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 823..1818  
; OTHER INFORMATION: /product= "arabinoxylan degrading  
; OTHER INFORMATION: enzyme" /gene= "axda"  
; OTHER INFORMATION: /standard\_name= "arabinoxylan degrading enzyme"  
; FEATURE:  
; NAME/KEY: sig\_peptide  
; LOCATION: 823..901  
; FEATURE:

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; NAME/KEY: mat_peptide
; LOCATION: 901..1818
US-08-637-763B-7

Query Match      48.1%; Score 17.8; DB 2; Length 2859;
Best Local Similarity 67.6%; Pred. No. 59;
Matches 25; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

QY 1 gcgcggatccatttcttagcataacggaagtc 37
    ||| ||||| ||||| || || ||||| ||
Db 2551 GCCTGGATCCCATTTGCTGTCGACAGCCTGGAACITC 2587

RESULT 9
US-09-170-354-7
; Sequence 7, Application US/09170354
; Patent No. 6066356
; GENERAL INFORMATION:
; APPLICANT: VAN DER WOUW, Monique J.A. et al
; TITLE OF INVENTION: ARABINOXYLAN DEGRADING ENZYME
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Morrison & Foerster
; STREET: 2000 Pennsylvania Avenue, NW
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20006-1812
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/170.354
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/637,763
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Murashige, Kate H.
; REGISTRATION NUMBER: 29,959
; REFERENCE/DOCKET NUMBER: 4615-0066.00
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 887-1500
; TELEFAX: (202) 887-0763
; TELEX: 90-4030 MRSNFOERSWSH
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2859 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Aspergillus niger var. tubigenis
; STRAIN: DS16813
; FEATURE:
; NAME/KEY: CAAT_signal
; LOCATION: 651..655
; FEATURE:
; NAME/KEY: TATA_signal
; LOCATION: 713..720
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 823..1818
; OTHER INFORMATION: /product= "arabinoxylan degrading
; OTHER INFORMATION: enzyme"
; OTHER INFORMATION: /gene= "axda"
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; OTHER INFORMATION: /standard_name= "arabinoxylan degrading enzyme"
; FEATURE:
; NAME/KEY: sig_peptide
; LOCATION: 823..901
; FEATURE:
; NAME/KEY: mat_peptide
; LOCATION: 901..1818
US-09-170-354-7

Query Match      48.1%; Score 17.8; DB 3; Length 2859;
Best Local Similarity 67.6%; Pred. No. 59;
Matches 25; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

QY 1 gcgcggatccatttcttagcataacggaagtc 37
    ||| ||||| ||||| || || ||||| ||
Db 2551 GCCTGGATCCCATTTGCTGTCGACAGCCTGGAACITC 2587

RESULT 10
US-09-522-666-1/c
; Sequence 1, Application US/09522666
; Patent No. 6333167
; GENERAL INFORMATION:
; APPLICANT: Shuey, David
; APPLICANT: Quinet, Elaine
; TITLE OF INVENTION: Methods and Reagents for Identifying Inhibitors of
; TITLE OF INVENTION: Proteolysis of Membrane-Associated Proteins
; FILE REFERENCE: 6-00
; CURRENT APPLICATION NUMBER: US/09/522,666
; CURRENT FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 1
; LENGTH: 3092
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence:contains
; OTHER INFORMATION: APF-Laci fusion protein
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (12)..(3083)
US-09-522-666-1

Query Match      47.6%; Score 17.6; DB 4; Length 3092;
Best Local Similarity 71.9%; Pred. No. 74;
Matches 23; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 4 ccggatccatttcttagcataacggaagt 35
    ||||| ||||| ||||| ||||| ||||| |||||
Db 1972 CGGATCCTTTCTTCTTCAGCATCACCAGGT 1941

RESULT 11
US-08-321-478-2
; Sequence 2, Application US/08321478
; Patent No. 5527677
; GENERAL INFORMATION:
; APPLICANT: DEGUCHI, Takeo
; APPLICANT: KINOSHITA, Moritoshi
; APPLICANT: KATSURAGI, Kiyonori
; APPLICANT: SHIN, Sadahito
; TITLE OF INVENTION: HUMAN ARYLAMINE N-ACETYLTRANSFERASE
; TITLE OF INVENTION: GENES
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Sughrue, Mion, Zinn, Macpeak & Seas
; STREET: 2100 Pennsylvania Avenue, N.W.
; CITY: Washington
; STATE: D.C.
; COUNTRY: United States
```

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; ZIP: 20037-3202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/321,478
; FILING DATE: 11-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/038,667
; FILING DATE: 23-MAR-1993
; APPLICATION NUMBER: JP 64669/1992
; FILING DATE: 23-MAR-1992
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 293-7060
; TELEFAX: (202) 293-7860
; TELEX: 6491103
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6464 base pairs
; TYPE: nucleic acid
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 723..1595
; FEATURE:
; NAME/KEY: exon
; LOCATION: 717..1936
; FEATURE:
; NAME/KEY: polyA_signal
; LOCATION: 1794..1799
; FEATURE:
; NAME/KEY: polyA_signal
; LOCATION: 1800..1805
; US-08-321-478-2

Query Match 47.0%; Score 17.4; DB 1; Length 6464;
Best Local Similarity 77.8%; Pred. No. 1.1e-02;
Matches 21; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 3 gcggatccatttccttagcataac 29
||| ||| ||| ||| ||| ||| ||| |||
Db 5386 GCTGGAGGCCATTATCCTTAGCAAAACC 5412

RESULT 12
US-08-321-478-4
; Sequence 4, Application US/08321478
; Patent No. 5527677
; GENERAL INFORMATION:
; APPLICANT: DEGUCHI, Takeo
; APPLICANT: KINOSHITA, Moritoshi
; APPLICANT: KATSURAGI, Kiyonori
; APPLICANT: SHIN, Sadahito
; TITLE OF INVENTION: HUMAN ARYLAMINE N-ACETYLTRANSFERASE
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Sughrue, Mion, Zinn, Macpeak & Seas
; STREET: 2100 Pennsylvania Avenue, N.W.
; CITY: Washington
; STATE: D.C.
; COUNTRY: United States
; ZIP: 20037-3202
; COMPUTER READABLE FORM:
; OPERATING SYSTEM: PC-DOS/MS-DOS
; COMPUTER: IBM PC compatible
; MEDIUM TYPE: Floppy disk
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/321,478
; FILING DATE: 11-OCT-1994
; CLASSIFICATION: 435
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; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/321,478
; FILING DATE: 11-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/038,667
; FILING DATE: 23-MAR-1993
; APPLICATION NUMBER: JP 64669/1992
; FILING DATE: 23-MAR-1992
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 293-7060
; TELEFAX: (202) 293-7860
; TELEX: 6491103
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6464 base pairs
; TYPE: nucleic acid
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 723..1595
; FEATURE:
; NAME/KEY: exon
; LOCATION: 717..1936
; FEATURE:
; NAME/KEY: polyA_signal
; LOCATION: 1794..1799
; FEATURE:
; NAME/KEY: polyA_signal
; LOCATION: 1800..1805
; US-08-321-478-4

Query Match 47.0%; Score 17.4; DB 1; Length 6464;
Best Local Similarity 77.8%; Pred. No. 1.1e-02;
Matches 21; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 3 gcggatccatttccttagcataac 29
||| ||| ||| ||| ||| ||| ||| |||
Db 5386 GCTGGAGGCCATTATCCTTAGCAAAACC 5412

RESULT 13
US-08-321-478-6
; Sequence 6, Application US/08321478
; Patent No. 5527677
; GENERAL INFORMATION:
; APPLICANT: DEGUCHI, Takeo
; APPLICANT: KINOSHITA, Moritoshi
; APPLICANT: KATSURAGI, Kiyonori
; APPLICANT: SHIN, Sadahito
; TITLE OF INVENTION: HUMAN ARYLAMINE N-ACETYLTRANSFERASE
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Sughrue, Mion, Zinn, Macpeak & Seas
; STREET: 2100 Pennsylvania Avenue, N.W.
; CITY: Washington
; STATE: D.C.
; COUNTRY: United States
; ZIP: 20037-3202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/321,478
; FILING DATE: 11-OCT-1994
; CLASSIFICATION: 435
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; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10754 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-966-958-1

Query Match 47.0%; Score 17.4; DB 2; Length 10754;
Best Local Similarity 77.8%; Pred. No. 1.2e+02;
Matches 21; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Qy 1 ggcgcggatccatttctcttagcata 27
      ||||| ||||| ||||| ||||| ||
Db 3992 GAGCTGGAGGCCATTATCCTTAGCAA 3966

RESULT 15
US-09-215-817-1/c
; Sequence 1, Application US/09215817
; Patent No. 5968786
; GENERAL INFORMATION:
; APPLICANT: Dunn, John
; APPLICANT: Randesi, Matthew
; TITLE OF INVENTION: METHODS FOR INTRODUCING UNIDIRECTIONAL
; TITLE OF INVENTION: DELETIONS
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brookhaven National Laboratory
; STREET: P.O. Box 5000
; CITY: Upton
; STATE: New York
; COUNTRY: US
; ZIP: 11973
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/215,817
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/966,958
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Bogosian, Margaret
; REGISTRATION NUMBER: 25,324
; REFERENCE/DOCKET NUMBER: AU197-14
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 344-3341
; TELEFAX: (516) 344-3729
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10754 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-09-215-817-1

Query Match 47.0%; Score 17.4; DB 2; Length 10754;
Best Local Similarity 77.8%; Pred. No. 1.2e+02;
Matches 21; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Qy 1 ggcgcggatccatttctcttagcata 27
      ||||| ||||| ||||| ||||| ||
Db 3992 GAGCTGGAGGCCATTATCCTTAGCAA 3966

```

Thu Aug 1 08:36:34 2002

us-09-824-567-4.rni

Page 7

Search completed: July 31, 2002, 19:33:02  
Job time: 8478 sec

Result	No.	Score	Query %			DB	ID	Description
			Match	Length	Match			
1	37	100.0	37	22	AAD20240	Chlamydia pneumoniae		
C	25.6	69.2	1799	22	AAD20238	Chlamydia pneumoniae		
C	25.6	69.2	1230025	20	AXX91990	Nucleotide sequence		
C	20.8	56.2	273254	21	AC81914	Chlamydia pneumoniae		
5	20.8	56.2	1230025	20	AXX91990	Nucleotide sequence		
C	20.6	55.7	238	21	AC97319	Helicobacter pylori		
C	20.6	55.7	448	21	AC97248	Helicobacter pylori		
C	20.6	55.7	474	21	AC97239	Helicobacter pylori		
C	20.6	55.7	3636	23	AAS3717	Helicobacter pylori		

## SUMMARIES

particular humans -

particular humans -

xx PS Claim 41; Page 53; 88pp; English.

xx CC The present invention relates to novel Chlamydia pneumoniae ATP-binding

cc cassette protein and its corresponding gene. Sequences of the invention

cc are useful for detecting Chlamydia infection by assaying a body fluid

cc of a mammal with the components. They are also used as vaccines. ATP

cc binding cassette antibodies and vaccines of the invention are useful

cc for preventing or treating Chlamydia infection e.g. infection caused

cc by C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum in mammals,

cc such as humans. The nucleic acid molecules are useful for producing

cc ATP-binding cassettes, in the construction of vaccine vectors such

cc as poxviruses, which are further useful for preventing and/or treating

cc Chlamydia infection and in the construction of attenuated Chlamydia

cc strains that can over-express the nucleic acid molecules or express

cc it in a non-toxic, mutated form. The present DNA sequence is a 3' PCR

cc primer which is used for amplifying Chlamydia pneumoniae ATP-binding

cc cassette DNA.

xx SQ Sequence 37 BP; 8 A; 12 C; 8 G; 9 T; 0 other;

Query Match 100.0%; Score 37; DB 22; Length 37;

Best Local Similarity 100.0%; Pred. No. 3.7e-07;

Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 gcgcggatccatttccttagcatacgaagtc 37

Db 1 gcgcggatccatttccttagcatacgaagtc 37

RESULT 2

AAD20238/C

ID AAD20238 standard; DNA; 1799 BP.

XX AC AAD20238;

XX DT 15-JAN-2002 (first entry)

XX DE Chlamydia pneumoniae ATP-binding cassette gene.

XX KW ATP-binding cassette; antibiotic; vaccine; infection; therapy; poxvirus;

xx ds.

xx OS Chlamydia pneumoniae.

xx FH Key Location/Qualifiers

FT CDS 101..1699

FT /\*tag= a

FT /product= "ATP-binding cassette protein"

XX PN W0200174863-A2.

XX PD 11-OCT-2001.

XX PF 04-APR-2001; 2001WO-CA00455.

XX PR 04-APR-2000; 2000US-194464P.

XX PA (AVET ) AVENTIS PASTEUR LTD.

PI Mordin AD, Oomen RP, Wang J, Dunn P;

XX WPI; 2001-648549/74.

DR P-PSDB; AAEL2212.

PT Novel Chlamydia ATP-binding cassette and corresponding DNA molecule for

PT preventing, diagnosing and treating Chlamydia infections in mammals, in

xx particular humans -

xx PS Claim 2; Fig 1; 88pp; English.

xx CC The present invention relates to novel Chlamydia pneumoniae ATP-binding

cc cassette protein and its corresponding gene. Sequences of the invention

cc are useful for detecting Chlamydia infection by assaying a body fluid

cc of a mammal with the components. They are also used as vaccines. ATP

cc binding cassette antibodies and vaccines of the invention are useful

cc for preventing or treating Chlamydia infection e.g. infection caused

cc by C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum in mammals,

cc such as humans. The nucleic acid molecules are useful for producing

cc ATP-binding cassettes, in the construction of vaccine vectors such

cc as poxviruses, which are further useful for preventing and/or treating

cc Chlamydia infection and in the construction of attenuated Chlamydia

cc strains that can over-express the nucleic acid molecules or express

cc it in a non-toxic, mutated form. The present sequence is a gene encoding

cc Chlamydia pneumoniae ATP-binding cassette.

xx SQ Sequence 1799 BP; 560 A; 439 C; 294 G; 506 T; 0 other;

Query Match 69.2%; Score 25.6; DB 22; Length 1799;

Best Local Similarity 87.5%; Pred. No. 0.11;

Matches 28; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 6 ggatccatttccttagcatacgaagtc 37

Db 1703 GGTGCTAATTTTCCTTAGCATACGGAAGTCC 1672

RESULT 3

AAX91990/C

ID AAX91990 standard; DNA; 1230025 BP.

XX AC AAX91990;

XX DT 13-SEP-1999 (first entry)

XX DE Nucleotide sequence of the complete genome of Chlamydia pneumoniae.

XX KW Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;

xx sinusitis; purulent otitis media; erythema nodosum; pharyngitis;

xx vaccine; neutralising epitope; ss.

xx OS Chlamydia pneumoniae.

xx PN W09927105-A2.

xx PD 03-JUN-1999.

xx PF 20-NOV-1998; 98WO-IB01890.

xx PR 04-NOV-1998; 98US-0107078.

xx PR 21-NOV-1997; 97FR-0014673.

xx PA (GEST ) GENSET.

xx PI Griffais R;

xx DR WPI; 1999-357842/30.

xx PT Genome sequence of Chlamydia pneumoniae

xx PS Claim 1; Page 291-611; 1912pp; English.

xx CC The present sequence represents the complete genome of Chlamydia

cc pneumoniae, and encodes proteins AAY34584-Y35879. C. pneumoniae causes

cc respiratory disease such as pneumonia and bronchitis and is thought

cc to be a contributing factor in heart disease, sarcoidosis, sinusitis,

cc purulent otitis media, erythema nodosum or pharyngitis. The polypeptides

cc encoded by the open reading frames of the C. pneumoniae genome (see

cc AAY34584-Y35879) can be used in immunogenic compositions as vaccines.

cc Vectors containing C. pneumoniae nucleotide sequences can also be

cc used as immunogenic compositions, especially where the vector directs

cc the expression of a neutralising epitope of C. pneumoniae.

xx SQ Sequence 1230025 BP; 367213 A; 249833 C; 249013 G; 363589 T; 377 other;



DE  
XX  
KW

KW antibacterial; bait polypeptide; gastric ulcer; ds.  
 XX Helicobacter pylori.  
 OS WO200066722-A1.  
 PN 09-NOV-2000.  
 XX 14-APR-2000; 2000WO-IB00603.  
 PF 30-APR-1999; 99EP-0401066.  
 XX (HYBR-) HYBRIGENICS SA.  
 PA Legrain P, Selig L, Rain J;  
 XX WPI; 2000-687535/67.  
 DR P-PSDB; AAB52573.  
 XX A two-hybrid system for identifying compounds useful in the treatment  
 PT of e.g. gastric ulcers comprises producing a collection of recombinant  
 PT cell clones -  
 XX Example 5; Page 197; 267pp; English.  
 XX The present sequence encodes a bait polypeptide used in a Helicobacter  
 CC pylori two-hybrid screen to identify protein-protein interactions.  
 CC The method is used to identify a recombinant cell clone expressing a  
 CC prey polypeptide which is capable of interacting with the bait  
 CC polypeptide. The two hybrid system is useful for screening compounds  
 CC for antibacterial activity. It may be used in the treatment of gastric  
 CC ulcers. The polynucleotides are useful as amplification primers or  
 CC specific detection probes. The polypeptides, vectors or host cells can  
 CC be used as immunogens to produce mono- or polyclonal antibodies. The  
 CC polynucleotides, polypeptides, antibodies, vectors, host cells or  
 CC modulating agents can be used to produce a pharmaceutical composition.  
 XX Sequence 238 BP; 56 A; 40 C; 64 G; 68 T; 0 other;

Query Match 55.7%; Score 20.6; DB 21; Length 238;  
 Best Local Similarity 85.2%; Pred. No. 13;  
 Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 6 ggatccatttccttagcataacgga 32  
 DB 103 GAATGCCCATTTCTTAGCATAACGGA 77  
 RESULT 7  
 AAC97248/G  
 ID AAC97248 standard; DNA; 448 BP.  
 XX AAC97248;  
 AC 23-FEB-2001 (first entry)  
 DT Helicobacter pylori bait polypeptide nucleotide sequence #20.  
 DE Helicobacter pylori; two-hybrid system; protein-protein interaction;  
 XX antibacterial; bait polypeptide; gastric ulcer; ds.  
 KW Helicobacter pylori.  
 OS WO200066722-A1.  
 PN 09-NOV-2000.  
 XX 14-APR-2000; 2000WO-IB00603.  
 PF 30-APR-1999; 99EP-0401066.  
 XX (HYBR-) HYBRIGENICS SA.  
 PA Legrain P, Selig L, Rain J;  
 XX WPI; 2000-687535/67.  
 DR P-PSDB; AAB52493.  
 XX A two-hybrid system for identifying compounds useful in the treatment  
 PT of e.g. gastric ulcers comprises producing a collection of recombinant  
 PT cell clones -  
 XX Example 5; Page 92-93; 267pp; English.  
 PS The present sequence encodes a bait polypeptide used in a Helicobacter  
 CC

XX Legrain P, Selig L, Rain J;  
 PI WPI; 2000-687535/67.  
 XX P-PSDB; AAB52502.  
 DR A two-hybrid system for identifying compounds useful in the treatment  
 OS of e.g. gastric ulcers comprises producing a collection of recombinant  
 PT cell clones -  
 XX Example 5; Page 102-103; 267pp; English.  
 XX The present sequence encodes a bait polypeptide used in a Helicobacter  
 CC pylori two-hybrid screen to identify protein-protein interactions.  
 CC The method is used to identify a recombinant cell clone expressing a  
 CC prey polypeptide which is capable of interacting with the bait  
 CC polypeptide. The two hybrid system is useful for screening compounds  
 CC for antibacterial activity. It may be used in the treatment of gastric  
 CC ulcers. The polynucleotides are useful as amplification primers or  
 CC specific detection probes. The polypeptides, vectors or host cells can  
 CC be used as immunogens to produce mono- or polyclonal antibodies. The  
 CC polynucleotides, polypeptides, antibodies, vectors, host cells or  
 CC modulating agents can be used to produce a pharmaceutical composition.  
 XX Sequence 448 BP; 143 A; 81 C; 102 G; 122 T; 0 other;  
 SQ Query Match 55.7%; Score 20.6; DB 21; Length 448;  
 Best Local Similarity 85.2%; Pred. No. 15;  
 Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 6 ggatccatttccttagcataacgga 32  
 DB 352 GAATGCCCATTTCTTAGCATAACGGA 326

RESULT 8  
 AAC97239/C  
 ID AAC97239 standard; DNA; 474 BP.  
 XX AAC97239;  
 AC 23-FEB-2001 (first entry)  
 DT Helicobacter pylori bait polypeptide nucleotide sequence #11.  
 DE Helicobacter pylori; two-hybrid system; protein-protein interaction;  
 XX antibacterial; bait polypeptide; gastric ulcer; ds.  
 KW Helicobacter pylori.  
 OS WO200066722-A1.  
 PN 09-NOV-2000.  
 XX 14-APR-2000; 2000WO-IB00603.  
 PF 30-APR-1999; 99EP-0401066.  
 XX (HYBR-) HYBRIGENICS SA.  
 PA Legrain P, Selig L, Rain J;  
 XX WPI; 2000-687535/67.  
 DR P-PSDB; AAB52493.  
 XX A two-hybrid system for identifying compounds useful in the treatment  
 PT of e.g. gastric ulcers comprises producing a collection of recombinant  
 PT cell clones -  
 XX Example 5; Page 92-93; 267pp; English.  
 PS The present sequence encodes a bait polypeptide used in a Helicobacter  
 CC

CC pylori two-hybrid screen to identify protein-protein interactions.  
 CC The method is used to identify a recombinant cell clone expressing a  
 CC prey polypeptide which is capable of interacting with the bait  
 CC polypeptide. The two hybrid system is useful for screening compounds  
 CC for antibacterial activity. It may be used in the treatment of gastric  
 CC ulcers. The polynucleotides are useful as amplification primers or  
 CC specific detection probes. The polypeptides, vectors or host cells can  
 CC be used as immunogens to produce mono- or polyclonal antibodies. The  
 CC polynucleotides, polypeptides, antibodies, vectors, host cells or  
 CC modulating agents can be used to produce a pharmaceutical composition.  
 XX  
 SQ Sequence 474 BP; 164 A; 91 C; 104 G; 115 T; 0 other;

Query Match 55.7%; Score 20.6; DB 21; Length 474;  
 Best Local Similarity 85.2%; Pred. No. 15;

Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 6 ggatccattttccttagcatacggga 32  
 | | | | |  
 Db 472 GAATGCCCATTTCTTAGCATACGGA 446

## RESULT 9

AA53717/c  
 ID AAS53717 standard; DNA; 3636 BP.

XX AC AAS53717;

XX DT 13-FEB-2002 (first entry)

XX DE Helicobacter pylori DNA for cellular proliferation protein #171.

XX KW Antisense; ds; prokaryotic cellular proliferation gene;

XX KW antibiotic; antibacterial; drug design.

XX OS Helicobacter pylori.

XX PN WO200170955-A2.

XX PD 27-SEP-2001.

XX PF 21-MAR-2001; 2001WO-US09180.

XX PR 21-MAR-2000; 2000US-191078P.

XX PR 23-MAY-2000; 2000US-206848P.

XX PR 26-MAY-2000; 2000US-207727P.

XX PR 23-OCT-2000; 2000US-242578P.

XX PR 27-NOV-2000; 2000US-253625P.

XX PR 22-DEC-2000; 2000US-257931P.

XX PR 16-FEB-2001; 2001US-269308P.

XX PA (ELIT-) ELITRA PHARM INC.

XX PI Haselbeck R, Ohlsen KL, Zyskind JW, Wall D, Trawick JD, Carr GJ;

XX PI Yamamoto RT, Xu HH;

XX DR WPI; 2001-611495/70.

XX DR P-PSDH; AAU35858.

XX PT New polynucleotides for the identification and development of  
 XX PT antibiotics, comprise sequences of antisense nucleic acids -

XX PS Claim 27; Seq ID No 7354; 511pp; English.

XX CC The invention relates to antisense inhibitors of genes essential to  
 XX CC prokaryotic cellular proliferation. their use in identifying the  
 XX CC genes themselves and the encoded proteins. The prokaryotes used are  
 XX CC Escherichia coli, Staphylococcus aureus, Salmonella typhi, Klebsiella  
 XX CC pneumoniae, Pseudomonas aeruginosa and Enterococcus faecalis. The  
 XX CC invention is also useful for the identification of potential new targets  
 XX CC for antibiotic development. The antisense nucleic acids can also be used

CC to identify proteins used in proliferation, to express these proteins,  
 CC and to obtain antibodies capable of binding to the expressed proteins.  
 CC The proteins can be used to screen compounds in rational drug discovery  
 CC programmes. The antisense nucleic acid sequence is also useful to screen  
 CC for homologous nucleic acids which are required for cell proliferation in  
 CC a wide variety of organisms. The present sequence encodes an  
 CC essential prokaryotic cellular proliferation protein.  
 CC Note: The sequence data for this patent did not form part  
 CC of the printed specification, but was obtained in electronic  
 CC format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.

XX SQ Sequence 3636 BP; 1184 A; 633 C; 873 G; 946 T; 0 other;

Query Match 55.7%; Score 20.6; DB 23; Length 3636;  
 Best Local Similarity 85.2%; Pred. No. 23;

Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 6 ggatccattttccttagcatacggga 32  
 | | | | |  
 Db 1141 GAATGCCCATTTCTTAGCATACGGA 1115

## RESULT 10

AA92471/c

ID AAX92471 standard; DNA; 20 BP.

XX AC AAX92471;

XX DT 13-SEP-1999 (first entry)

XX DE PCR primer used to amplify an ORF of Chlamydia pneumoniae.

XX KW Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;

XX KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;

XX KW vaccine; neutralising epitope; PCR primer; ss.

XX OS Synthetic.

XX OS Chlamydia pneumoniae.

XX PN WO927105-A2.

XX PD 03-JUN-1999.

XX PF 20-NOV-1998; 98WO-IB01890.

XX PR 04-NOV-1998; 98US-0107078.

XX PR 21-NOV-1997; 97FR-0014673.

XX PA (GEST ) GENSET.

XX PI Griffais R;

XX DR WPI; 1999-357842/30.

XX PT Genome sequence of Chlamydia pneumoniae

XX PS Page 1514; Disclosure; 1912pp; English.

XX CC AAX91991-X97517 represent PCR primers used to amplify open reading  
 XX CC frames and other nucleic acid sequences from the genome of  
 XX CC Chlamydia pneumoniae (see AAX91990). C. pneumoniae causes respiratory  
 XX CC disease such as pneumonia and bronchitis and is thought to be a  
 XX CC contributing factor in heart disease, sarcoidosis, sinusitis, purulent  
 XX CC otitis media, erythema nodosum or pharyngitis. The polypeptides encoded  
 XX CC by the open reading frames of the C. pneumoniae genome (see AAX34584-  
 XX CC AAX35879) can be used in immunogenic compositions as vaccines. Vectors  
 XX CC containing C. pneumoniae nucleotide sequences can also be used as  
 XX CC immunogenic compositions, especially where the vector directs the  
 XX CC expression of a neutralising epitope of C. pneumoniae.

XX SQ Sequence 20 BP; 4 A; 4 C; 6 G; 6 T; 0 other;

Query Match 54.1%; Score 20; DB 20; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 14;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 18 ccttagcataacgaagtc 37  
 Db 20 CCTTAGCATACGGAAGTCC 1

RESULT 11  
 AAV38294/C  
 ID AAV38294 standard; cDNA; 362 BP.  
 XX  
 AC AAV38294;  
 XX  
 DT 12-OCT-1998 (first entry)  
 XX  
 DE Human C-C chemokine DGWCC cDNA.  
 XX  
 KW DGWCC; DNAX groin wound expressed CC chemokine; cytokine; human;  
 KW immune system; cancer; cell proliferation; therapy; diagnosis; ss.  
 XX  
 OS Homo sapiens.

Key Location/Qualifiers  
 CDS 1..339  
 FT /\*tag= a  
 FT sig\_peptide 1..72  
 FT /\*tag= b  
 FT mat\_peptide 73..336  
 FT /\*tag= c

PN W09823750-A2.  
 XX  
 PD 04-JUN-1998.  
 XX  
 PF 26-NOV-1997; 97WO-US21092.  
 XX  
 PR 05-DEC-1996; 96US-0761071.  
 PR 27-NOV-1996; 96US-0031805.  
 XX  
 PA (SCHE ) SCHERING CORP.  
 XX  
 PI Hedrick JA, Morales J, Vicari A, Zlotnik A;  
 XX  
 DR WPI; 1998-322730/28.  
 DR P-PSDB; AAW60650.

PT Dvic-1 and DGWCC chemokines - useful for developing products for  
 PT treating abnormal physiology or development, e.g. cancerous or  
 PT degenerative conditions  
 XX  
 PS Disclosure; Page 62; 71pp; English.

XX This cDNA sequence codes for novel human DNAX groin wound expressed  
 CC chemokine (DGWCC) (see AAW60650). DGWCC cDNA can be obtained from  
 CC e.g. skin, epithelial or wound healing libraries by PCR  
 CC amplification or by hybridisation. Also disclosed is novel human  
 CC DNAX vic-1 (DVic-1) (see AAW60649), as well as expression vectors and  
 CC host cells. DGWCC and DVic-1 play a role in the regulation or  
 CC development of neuronal or haematopoietic cells, e.g. lymphoid  
 CC cells, which affect immunological responses. They can be used in  
 CC the treatment of conditions associated with abnormal physiology or  
 CC development, including abnormal proliferation, e.g. cancerous  
 CC conditions or degenerative conditions. Abnormal proliferation,  
 CC regeneration, therapeutic treatment, and atrophy may be modulated by  
 CC appropriate therapeutic treatment using products of the invention.  
 CC The products can also be used for detection, diagnosis and drug  
 CC screening.

XX Sequence 362 BP; 91 A; 116 C; 85 G; 70 T; 0 other;  
 SQ

Query Match 54.1%; Score 20; DB 19; Length 362;  
 Best Local Similarity 72.2%; Pred. No. 27;  
 Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 ggcgcggatccatttccttagcataacgaagtc 36  
 Db 344 GGCCTTCAGCCCATTTTCCTTAGCATCCCAAAATTC 309

RESULT 12  
 AAA47548/C  
 ID AAA47548 standard; DNA; 362 BP.  
 XX  
 AC AAA47548;  
 XX  
 DT 20-OCT-2000 (first entry)  
 XX  
 DE Primate CTACK nucleotide sequence.  
 XX  
 KW Cutaneous T-cell attracting chemokine; CTACK; skin; cell movement;  
 KW migration; vasoactive intestinal contractor; Vic; GPR2; agonist;  
 KW antagonist; antibody; immunological condition; mutein; ds.  
 XX  
 OS Homo sapiens.

Key Location/Qualifiers  
 CDS 1..339  
 FT /\*tag= a  
 FT /product= CTACK 1..72  
 FT sig\_peptide 1..72  
 FT /\*tag= b  
 FT mat\_peptide 73..336  
 FT /\*tag= c

PN W0200038713-A1.  
 XX  
 PD 06-JUL-2000.  
 XX  
 PF 23-DEC-1999; 99WO-US30819.  
 XX  
 PR 24-DEC-1998; 98US-0113858.  
 PR 27-MAY-1999; 98US-0322580.  
 XX  
 PA (SCHE ) SCHERING CORP.  
 XX  
 PI Wang W, Oldham ER, Soto H, Lui Y, Hudak SA, Homey B, Morales JM;  
 PI Kellermann S, McEvoy LM, Zlotnik A;  
 XX  
 DR WPI; 2000-465633/40.  
 DR P-PSDB; AAB01453.

PT Modulating cell movement within the skin, useful for treating  
 PT immunological skin conditions or diseases comprises administering T  
 PT cell-attracting chemokine or vasoactive intestinal contractor chemokine  
 PT agonists or antagonists

XX Example 3; Page 73; 79pp; English.

XX Modulating movement of a cell within or to the skin of a mammal can  
 CC be achieved by administering an antagonist or agonist of cutaneous T  
 CC cell-attracting chemokine (CTACK) or vasoactive intestinal contractor  
 CC (Vic) chemokine. The antagonist is selected from a mutein of natural  
 CC CTACK or Vic, an antibody which neutralises CTACK or Vic or an  
 CC antibody which block GPR2 ligand binding. The CTACK or Vic agonists  
 CC or antagonists are useful for treating medical conditions or diseases  
 CC associated with immunological conditions of the skin.

XX Sequence 362 BP; 91 A; 116 C; 85 G; 70 T; 0 other;  
 SQ

Query Match 54.1%; Score 20; DB 21; Length 362;



PA (IPKP-) IPK INST PFLANZENGENETIK & KULTURPFLANZE.  
 XX Koerner M, Berndt E, Fritsche K, Feussner I;  
 XX WPI; 2002-098656/14.  
 XX New polynucleotide, useful for producing transgenic food plants with  
 PT altered contents of polyene fatty acids, comprises recombinant nucleic  
 PT acid encoding plant acylhydrolase -  
 XX  
 XX Claim 5; Page 12; 25pp; German.  
 XX  
 CC The present invention provides coding sequences and proteins from  
 CC Arabidopsis thaliana which act as acylhydrolase enzymes. These are  
 CC designated triacylglycerol lipase (TAG-lipase) 1,2 and 3. The sequences  
 CC are used to produce transgenic plants, which are useful in human or  
 CC animal nutrition and have altered content of (oxygenated) polyene fatty  
 CC acids (PEFA) in the seed oil. They can also be used to identify and  
 CC isolate other TAG-lipase encoding sequences from genomic/cDNA libraries.  
 CC The present sequence is the TAG-lipase 1 coding sequence of the  
 CC invention.  
 CC Note: The present sequence is stated in the specification as encoding the  
 CC protein shown in SEQ ID NO: 2 (ABB04482). However, this is not the case.  
 CC  
 XX Sequence 1246 BP; 324 A; 263 C; 296 G; 363 T; 0 other;

Query Match 53.0%; Score 19.6; DB 24; Length 1246;  
 Best Local Similarity 84.6%; Pred. No. 52;  
 Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 9 tccattttctcttagcataacggaag 34  
 | | ||||| ||||| |||||  
 Db 838 ttcaattttctcttagcataacggaag 863

Search completed: July 31, 2002, 20:59:28  
 Job time: 7759 sec

GenCore version 4.5  
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 31, 2002, 19:31:22 ; Search time 1882.78 seconds  
(without alignments)  
411.244 Million cell updates/sec

Title: US-09-824-567-4

Perfect score: 37

Sequence: 1 ggcgcggatccatttcttagcatacgaagtc 37

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:

1: gb.ba.\*

2: gb.btg.\*

3: gb.in.\*

4: gb.om.\*

5: gb.ov.\*

6: gb.pat.\*

7: gb.ph.\*

8: gb.pl.\*

9: gb.pr.\*

10: gb.ro.\*

11: gb.sts.\*

12: gb.sy.\*

13: gb.un.\*

14: gb.vi.\*

15: em.ba.\*

16: em.fun.\*

17: em.hum.\*

18: em.in.\*

19: em.mu.\*

20: em.om.\*

21: em.or.\*

22: em.ov.\*

23: em.pat.\*

24: em.ph.\*

25: em.pl.\*

26: em.ro.\*

27: em.sts.\*

28: em.un.\*

29: em.vi.\*

30: em.htg.hum.\*

31: em.htg.inv.\*

32: em.htg.other.\*

33: em.htgo.inv.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
-----					

1	37	100.0	37	6	AX268344	Sequence
2	25.6	69.2	1799	6	AX268341	Sequence
3	25.6	69.2	11648	1	AE001606	Chlamydia
4	25.6	69.2	11764	1	AE002216	Chlamydia
5	25.6	69.2	299650	1	AP002545	Chlamydia
6	25	67.6	1599	6	AX349501	Sequence
7	21.6	58.4	166919	2	AL591675	Mus muscu
8	21.6	58.4	187334	2	AL626766	Mus muscu
9	21.6	58.4	226999	2	AC105488	Rattus no
10	21.6	58.4	239486	2	AC097752	Rattus no
11	21.4	57.8	151321	9	AC069483	Homo sapi
12	21.4	57.8	170257	9	AC078909	Homo sapi
13	21.2	57.3	70148	9	AL357061	Homo sapi
14	21.2	57.3	107967	9	AL353701	Human DNA
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SOURCE		synthetic construct.					
ORGANISM		artificial construct.					
REFERENCE		1 (sites)					
AUTHORS		Murkin,A.D., Oomen,R.P., Wang,J. and Dunn,P.					
TITLE		Chlamydia antigens and corresponding dna fragments and uses thereof					
JOURNAL		Patent: WO 0174863-A 4 11-OCT-2001;					
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Genome sequences of Chlamydia trachomatis MoPn and Chlamydia pneumoniae AR39
Nucleic Acids Res. 28 (6), 1397-1406 (2000)

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TITLE Chlamydomophila pneumoniae AR39
JOURNAL Nucleic Acids Res. 28 (6), 1397-1406 (2000)
MEDLINE 20150255
PUBMED 10684935
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AUTHORS Read,T.D., Brunham,R.C., Shen,C., Gill,S.R., Heidelberg,J.F., White,O., Hickey,E.K., Peterson,J., Umayam,L.A., Utterback,T., Berry,K., Bass,S., Linher,K., Weidman,J., Khouri,H., Craven,B., Bowman,C., Dodson,R., Gwinn,M., Nelson,W., DeBoy,R., Kolonay,J., McClarty,G., Salzberg,S.L., Eisen,J. and Fraser,C.M.
Direct Submission

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TITLE Chlamydomophila pneumoniae AR39
JOURNAL Submitted (01-MAR-2000) The Institute for Genomic Research, 9712 Medical Center Dr, Rockville, MD 20850, USA
COMMENT On Jun 1, 2000 this sequence version replaced gi:7189484.
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Db 105 GGTGCTAAATTTCCCTAGCATACGAAGTCC 136

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 AP002545.2 GI:9956082  
 ORGANISM Chlamydothila pneumoniae J138 (strain:J138) DNA.  
 Chlamydothila pneumoniae J138  
 Bacteria; Chlamydiales; Chlamydiaceae; Chlamydothila.  
 1 (sites)  
 Shirai,M., Hirakawa,H., Kimoto,M., Tabuchi,M., Kishi,F., Ouchi,K.,  
 Shiba,T., Ishii,K., Hattori,M., Kubara,S. and Nakazawa,T.  
 Comparison of whole genome sequences of Chlamydia pneumoniae J138  
 from Japan and CNL029 from USA  
 Nucleic Acids Res. 28 (12), 2311-2314 (2000)  
 20330349  
 2 (bases 1 to 299650)  
 Shirai,M.  
 Direct Submission  
 Submitted (04-JUL-2000) Mutsunori Shirai, Yamaguchi University  
 School of Medicine, Department of Microbiology, 1-1-1  
 Minamikoogushi, Ube, Yamaguchi 755-8505, Japan  
 (E-mail:mshirai@po.cc.yamaguchi-u.ac.jp, Tel:81-836-22-2227,  
 Fax:81-836-22-2415)  
 On Aug 31, 2000 this sequence version replaced gi:6172286  
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 AB033780-AB033781, AB033792-AB033799: Submitted (25-Oct-1999)  
 AB038345-AB038347: Submitted (14-Feb-2000)  
 AB036071-AB036078: Submitted (18-Dec-2000).  
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RESULT 9  

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LOCUS  

DEFINITION  

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SEQUENCE, 13 unordered pieces.  

AC105488  

AC105488.1 GI:18092710  

HTG; HTGS-PHASE1; HTGS_DRAFT; HTGS_FULLTOP.  

KEYWORDS  

Norway rat.  

SOURCE  

ORGANISM  

Rattus norvegicus  

Eukaryota; Metazoa;  

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  

Rattus.  

REFERENCE  

1 (bases 1 to 226999)  

Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,  

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Query Match 58.4%; Score 21.6; DB 2; Length 226999;  
 Best Local Similarity 75.0%; Pred. No. 1.3e+02;  
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RESULT 10  
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 SOURCE Norway rat.  
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 1 (bases 1 to 239486)  
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 Weinstein,G. and Gibbs,R.  
 Direct Submission  
 unpublished  
 2 (bases 1 to 239486)  
 Worley,K.C.  
 Direct Submission  
 TITLE  
 JOURNAL  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL  
 COMMENT  
 Submitted (23-OCT-2001) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 On Dec 20, 2001 this sequence version replaced gi:16327457.  
 ----- Genome Center  
 Center: Baylor College of Medicine

Center code: BCM  
 Web site: <http://www.hgsc.bcm.tmc.edu/>  
 Contact: hgsc-help@bcm.tmc.edu  
 ----- Project Information  
 Center project name: GFAW  
 Center clone name: CH230-75H6  
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 Consensus quality: 228942 bases at least Q20  
 Estimated insert size: 228246; sum-of-contigs estimation  
 Quality coverage: 9x in Q20 bases; agarose-fp estimation  
 Quality coverage: 4.5x in Q20 bases; sum-of-contigs estimation  
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\* NOTE: Estimated insert size may differ from sequence length  
 (see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
 \* NOTE: This is a 'working draft' sequence. It currently  
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 is not known and their order in this sequence record is  
 arbitrary. Gaps between the contigs are represented as  
 runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 as soon as it is available and the accession number will  
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 \* 62034 62133: gap of unknown length  
 \* 62134 73808: contig of 11675 bp in length  
 \* 73809 73908: gap of unknown length  
 \* 73909 94892: contig of 20984 bp in length  
 \* 94893 94992: gap of unknown length  
 \* 94993 111560: contig of 16568 bp in length  
 \* 111561 111561: gap of unknown length  
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 \* 124504 124504: gap of unknown length  
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 \* 141810 141909: gap of unknown length  
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* 215001 215002: contig of 1332 bp in length
* 215003 215004: gap of unknown length
* 215005 215006: contig of 1370 bp in length
* 215007 215008: gap of unknown length
* 215009 215010: contig of 1148 bp in length
* 215011 215012: gap of unknown length
* 215013 215014: contig of 1064 bp in length.

```

## FEATURES

```

Source
1..239486
/organism="Rattus norvegicus"
/db_xref="taxon:10116"
/chromosome="Rf1"
/clone="CH230-75H6"
64295 a 53502 c 53422 g 64642 t 3625 others

```

```

BASE COUNT 64295 a 53502 c 53422 g 64642 t 3625 others
ORIGIN

```

## Query Match

```

Best Local Similarity 58.4%; Score 21.6; DB 2; Length 239486;
Matches 27; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

```

```

QY 2 cgcgcgacccatttccttagcataacgaagtc 37

```

```

Db 162498 CGCTGGTCCTGTTTCCCTTCTGTCGACAGAGTCC 162463

```

## RESULT 11

```

AC069483
LOCUS AC069483 151321 bp DNA linear PRI 25-NOV-2001
DEFINITION Homo sapiens, clone RP11-29M5, complete sequence.
ACCESSION AC069483
VERSION AC069483.6 GI:17064696
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 151321)
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens, clone RP11-29M5
Unpublished
2 (bases 1 to 151321)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
Boguslavsky,L., Bouckhgalter,B., Brown,A., Burkett,G.,
Campopiano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
Collamore,A., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Grant,G., Hagos,B., Hearford,A., Horton,L.,
Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,I., Karatas,A.,
Klein,J., LaRocque,K., Lamazares,R., Landers,T., Lehoczy,J.,
Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N.,
McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,

```

## FEATURES

## Source

```

1..151321
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-29M5"
/clone_lib="RPC1-11 Human Male BAC"
920..1158
/rpt_family="MIR"
/rpt_family="AT_rich"
1519..1541
1829..2044
/rpt_family="AluJb"
2057..2351
/rpt_family="AluJo"
complement(2608..2684)
/rpt_family="MIR"
complement(5550..5748)
/rpt_family="MIR"
5809..5935
/rpt_family="FLAM_A"
7165..7216
/rpt_family="AT_rich"

```

## repeat\_region

## repeat\_region

## repeat\_region

## repeat\_region

## repeat\_region

## repeat\_region

## repeat\_region

## repeat\_region

Meldrim,J., Meneus,L., Mihova,T., Miranda,C., Mlenga,V., Morrow,J.,  
 Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,  
 O'Neill,D., Olivari,T.M., Oliver,J., Peterson,K., Pierre,N.,  
 Pisani,D., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,  
 Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,  
 Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,  
 Tesfaye,S., Theodore,J., Tirrell,A., Travers,M., Trigilio,J.,  
 Vassiliev,H., Viel,R., Vo.A., Wilson,B., Wu.X., Wyman,D., Ye,W.J.,  
 Young,G., Zainoun,J., Zimmer,A. and Zody,M.

## TITLE

## JOURNAL

Submitted (01-JUN-2000) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA

## REFERENCE

## AUTHORS

3 (bases 1 to 151321)  
 Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,  
 Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Bouckhgalter,B.,  
 Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,  
 Choepel,Y., Colangelo,M., Collins,S., Collamore,A., Cooke,A.,  
 Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,  
 Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,  
 Ginde,S., Goyette,M., Graham,L., Hulme,W., Iliev,I., Johnson,R.,  
 Hagos,B., Hearford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,  
 Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,  
 Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,  
 MacLean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,  
 McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrim,J.,  
 Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,  
 Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D.,  
 Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,  
 Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupbach,R.,  
 Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,  
 Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,  
 Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,  
 Viel,R., Vo.A., Wilson,B., Wu.X., Wyman,D., Ye,W.J., Young,G.,  
 Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

## TITLE

## JOURNAL

Submitted (25-NOV-2001) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA

On Nov 25, 2001 this sequence version replaced gi:16756289.

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L10441

Center clone name: 29\_M\_5

## FEATURES

## Source

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1..151321
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-29M5"
/clone_lib="RPC1-11 Human Male BAC"
920..1158
/rpt_family="MIR"
/rpt_family="AT_rich"
1519..1541
1829..2044
/rpt_family="AluJb"
2057..2351
/rpt_family="AluJo"
complement(2608..2684)
/rpt_family="MIR"
complement(5550..5748)
/rpt_family="MIR"
5809..5935
/rpt_family="FLAM_A"
7165..7216
/rpt_family="AT_rich"

```



```

repeat_region      8907..9195
                    /rpt_family="AluJo"
                    complement(9196..9478)
repeat_region      9488..9510
                    /rpt_family="AluSx"
repeat_region      9768..9789
                    /rpt_family="AT-rich"
repeat_region      11461..11483
                    /rpt_family="AT-rich"
repeat_region      complement(13745..13867)
                    /rpt_family="L2"
repeat_region      14034..14220
                    /rpt_family="MIR"
repeat_region      14417..14722
                    /rpt_family="AluJb"
repeat_region      complement(14991..15055)
                    /rpt_family="MER116"
repeat_region      16845..16866
                    /rpt_family="MIR3"
repeat_region      complement(17501..17900)
                    /rpt_family="L1ME1"
repeat_region      complement(18075..18202)
                    /rpt_family="MIR"
repeat_region      18203..18303
                    /rpt_family="HY3"
repeat_region      19347..19655
                    /rpt_family="AluSx"
repeat_region      complement(19674..19845)
                    /rpt_family="L1MB7"
repeat_region      22277..22296
                    /rpt_family="MIR"
repeat_region      23628..23654
                    /rpt_family="MIR"
repeat_region      complement(24237..24594)
                    /rpt_family="L2"
repeat_region      24562..25061
                    /rpt_family="MIR"
repeat_region      25148..25452
                    /rpt_family="AluSc"
repeat_region      26584..26712
                    /rpt_family="MIR"
repeat_region      complement(26967..27238)
                    /rpt_family="AluJo"
repeat_region      complement(27810..27952)
                    /rpt_family="MER5A"
repeat_region      complement(28204..28348)
                    /rpt_family="MER5A"
repeat_region      29028..29062
                    /rpt_family="CA)n"
repeat_region      complement(34139..34273)
                    /rpt_family="MIR3"
repeat_region      36575..36597
                    /rpt_family="MIR"
repeat_region      37037..37270
                    /rpt_family="CA)n"
repeat_region      37273..37317
                    /rpt_family="AluJo"
repeat_region      37685..37711
                    /rpt_family="CA)n"
repeat_region      37718..37765
                    /rpt_family="CA)n"
repeat_region      37876..38051
                    /rpt_family="GA-rich"
repeat_region      38194..38492
                    /rpt_family="MIR"
repeat_region      38497..38753
                    /rpt_family="AluJo"
repeat_region      complement(39331..39251)
                    /rpt_family="AluSx"
repeat_region      /rpt_family="FLAM_C"
                    39305..39411
repeat_region      /rpt_family="MIR"
                    41050..41368

```

```

repeat_region      /rpt_family="L1MC1"
                    43352..43379
repeat_region      /rpt_family="AT-rich"
                    43720..43839
repeat_region      /rpt_family="MIR"
                    44936..45221
repeat_region      /rpt_family="AluSg"
                    45625..45873
repeat_region      /rpt_family="L1ME4A"
                    45963..46178
repeat_region      /rpt_family="L1ME4A"
                    46179..46490
repeat_region      /rpt_family="AluSg"
                    46491..46583
repeat_region      /rpt_family="L1ME4A"
                    46585..46783
repeat_region      /rpt_family="AluJo"
                    46903..47289
repeat_region      /rpt_family="L1ME4A"
                    47404..47427
repeat_region      /rpt_family="AT-rich"
                    complement(47968..48272)
repeat_region      /rpt_family="MLT1B"
                    49874..49970
repeat_region      /rpt_family="MIR"
                    50992..51146
repeat_region      complement(51519..51809)

```

```

Query Match      57.8%; Score 21.4; DB 9; Length 151321;
Best Local Similarity 80.6%; Pred. No. 1.6e+02;
Matches 25; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

```

```

QY      6 ggatccatttccttagcatacaggaagtc 36
        || |||| |||| |||| |||| |||| |||| ||
Db      30449 GGGTCCTTTTCTTATCATACAGAGGC 30479

```

```

RESULT 12
AC078909      AC078909      170257 bp      DNA      linear      PRI 12-DEC-2001
LOCUS      Homo sapiens chromosome 15, clone RP11-128A17, complete sequence.
DEFINITION      AC078909
ACCESSION      AC078909
VERSION      AC078909.7 GI:17530779
KEYWORDS      HTG.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 170257)
AUTHORS      Birren, B., Linton, L., Nusbaum, C. and Lander, E.
TITLE      Homo sapiens chromosome 15, clone RP11-128A17
JOURNAL      Unpublished
REFERENCE      2 (bases 1 to 170257)
AUTHORS      Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
Anderson, S., Baldwin, J., Barna, N., Bastien, V., Beda, F.,
Boguslavskiy, L., Boukhalter, B., Brown, A., Burkett, G.,
Campopiano, A., Castle, A., Choepel, Y., Colangelo, M., Collins, S.,
Collamore, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S.,
Dodge, S., Domino, M., Doyle, M., Ferreira, P., FitzHugh, W., Gage, D.,
Galagan, J., Gardyna, S., Ginde, S., Goyette, M., Graham, L.,
Grand-pierre, N., Grant, G., Hagos, B., Heaford, A., Horton, L.,
Howland, J.C., Iliev, I., Johnson, R., Jones, C., Kann, L., Karatas, A.,
Klein, J., Laroque, K., Lamazares, R., Landers, T., Lehoczy, J.,
Levine, R., Lieu, C., Liu, G., Locke, K., Macdonald, P., Marquis, N.,
McCarthy, M., McEwan, P., McGuck, A., McKernan, K., McPheeters, R.,
Meldrum, J., Mepeus, L., Mihova, T., Miranda, C., Mienga, V., Morrow, J.,
Murphy, T., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell, P.,
O'Neil, D., Olivier, T.M., Oliver, J., Peterson, K., Pierre, N.,
Pisani, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D.,
Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B.,
Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,
Teschfaye, S., Theodore, J., Tirrell, A., Travers, M., Trigilio, J.,

```

TITLE  
JOURNAL  
REFERENCE  
AUTHORS

2  
3  
4  
5  
6  
7  
8  
9

TITLE  
JOURNAL

## REFERENCE AUTHORS

TYPE

## MENT

source

1

```

/db_xref="taxon:9606"
/chr="1"
/chromosome="15"
/clone="RP11-128A17"
/clone_lib="RPCI-II Hum
complement(1..99)
/rpt_family="MER18"
209..394
/rpt_family="AluSc"
1020..1068
/rpt_family="(TC)n"
1068..1104
/rpt_family="(TG)n"
2513..2533
/rpt_family="AT-rich"
complement(2616..2711)
/rpt_family="L2"
complement(2918..3086)
/rpt_family="MIR"
4316..4347
/rpt_family="(TA)n"
4348..4375
/rpt_family="(TA)n"
complement(4456..4494)
/rpt_family="L2"
complement(4563..4977)
/rpt_family="Tiger2"
5199..5488
/rpt_family="AluSq"
complement(5658..5877)
/rpt_family="THER1C"
complement(5877..6081)
/rpt_family="THER1C"
complement(6413..6734)
/rpt_family="AluJo"
6953..7269
/rpt_family="AluSx"
7286..7318
/rpt_family="(CAGG)n"
7498..7591
/rpt_family="MER45A"
complement(7592..8322)
/rpt_family="LPA10"
8323..8405
/rpt_family="MER45A"
complement(8434..8488)
/rpt_family="MER94"
8592..8886
/rpt_family="AluY"
10059..10237
/rpt_family="LIME"
11324..11367
/rpt_family="AT-rich"
complement(11666..11987)
/rpt_family="LTR16A"
complement(12098..12265)
/rpt_family="HERV16"
complement(12345..12443)
/rpt_family="HERV16"
12530..12737
/rpt_family="MLTIE"
complement(12811..13097)
/rpt_family="AluSx"
complement(13748..13907)
/rpt_family="HERV16"
complement(13908..14210)
/rpt_family="AluSx"
complement(14211..15723)
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/rpt_family="HERV16"

```

[illegible]

---

D., D., Ye, W. J.,  
 r for Genome  
 A  
 Allen, N.,  
 Kkhgalter, B.,  
 ro, B.,  
 ook, A.,  
 S.,  
 S.,  
 e, N.,  
 Johnson, R.,  
 G.,  
 S.,  
 aldrim, J.,  
 N.,  
 sil, D.,  
 ra, V.,  
 ogov, P.,  
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 ev, H.,  
 ough, G.,  
 for Genome  
 631.  
 Research

[illegible][illegible][illegible][illegible]



Rp11-203M2 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.  
 The true left end of clone Rp11-203M2 is at 1 in this sequence. The true left end of clone Rp11-738I14 is at 107868 in this sequence. The true right end of clone Rp11-479K20 is at 84404 in this sequence.

## FEATURES

Location/Qualifiers

source

1..107967  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /chromosome="9"  
 /clone="Rp11-203M2"  
 /clone\_lib="RPC1-11.1"

repeat\_region 178..724  
 /note="L2 repeat: matches 1850..2403 of consensus"  
 repeat\_region 864..904  
 /note="5S repeat: matches 2..42 of consensus"  
 repeat\_region 1407..1731  
 /note="AluY repeat: matches 1..306 of consensus"  
 repeat\_region 2351..2487  
 /note="AluSx repeat: matches 1..143 of consensus"  
 repeat\_region 2488..2801  
 /note="AluY repeat: matches 1..311 of consensus"  
 repeat\_region 2802..2989  
 /note="AluSx repeat: matches 143..311 of consensus"  
 repeat\_region 3322..3780  
 /note="L1MB6 repeat: matches 5706..6173 of consensus"  
 repeat\_region 3781..3888  
 /note="HAL1 repeat: matches 53..166 of consensus"  
 repeat\_region 3974..4146  
 /note="HAL1 repeat: matches 161..355 of consensus"  
 repeat\_region 4152..4296  
 /note="HAL1 repeat: matches 798..954 of consensus"  
 repeat\_region 4412..4732  
 /note="HAL1 repeat: matches 1177..1540 of consensus"  
 repeat\_region 4733..5042  
 /note="AluJb repeat: matches 1..308 of consensus"  
 repeat\_region 5043..5097  
 /note="HAL1 repeat: matches 1540..1588 of consensus"  
 repeat\_region 5568..5716  
 /note="L2 repeat: matches 2335..2516 of consensus"  
 repeat\_region 5772..5978  
 /note="L2 repeat: matches 2641..2749 of consensus"  
 repeat\_region 6188..6497  
 /note="AluY repeat: matches 1..311 of consensus"  
 repeat\_region 6732..6757  
 /note="L2 repeat: matches 2634..2656 of consensus"  
 repeat\_region 6758..7065  
 /note="AluSx repeat: matches 1..302 of consensus"  
 repeat\_region 7066..7124  
 /note="L2 repeat: matches 2656..2728 of consensus"  
 repeat\_region 7179..7351  
 /note="AluSg/x repeat: matches 127..299 of consensus"  
 repeat\_region 13752..14046  
 /note="AluJb repeat: matches 1..302 of consensus"  
 repeat\_region 14210..14520  
 /note="AluSx repeat: matches 1..312 of consensus"  
 repeat\_region 14522..14571  
 /note="25 copies 2 mer ac 92% conserved"  
 repeat\_region 14872..15071  
 /note="MER3 repeat: matches 1..201 of consensus"  
 repeat\_region 15292..15588  
 /note="AluSx repeat: matches 16..310 of consensus"  
 repeat\_region 16222..16529  
 /note="AluY repeat: matches 1..307 of consensus"  
 repeat\_region 16594..16755  
 /note="FRAM repeat: matches 0..161 of consensus"  
 repeat\_region 17073..17364  
 /note="AluSx repeat: matches 1..293 of consensus"  
 repeat\_region 17721..17894  
 /note="L1MC5 repeat: matches 7505..7669 of consensus"  
 repeat\_region 17895..18185  
 /note="AluSx repeat: matches 1..288 of consensus"

repeat\_region 18186..18399  
 /note="L1MC5 repeat: matches 7669..7913 of consensus"  
 repeat\_region 18879..19203  
 /note="L1MC4 repeat: matches 7091..7447 of consensus"  
 repeat\_region 19204..19503  
 /note="AluSg repeat: matches 1..299 of consensus"  
 repeat\_region 19504..19624  
 /note="L1MC4 repeat: matches 7447..7570 of consensus"  
 repeat\_region 19701..19843  
 /note="L1F5 repeat: matches 5316..5458 of consensus"  
 repeat\_region 19871..20175  
 /note="AluJo repeat: matches 1..301 of consensus"  
 repeat\_region 20315..20581  
 /note="AluJb repeat: matches 1..298 of consensus"  
 repeat\_region 20732..21027  
 /note="AluSg repeat: matches 1..294 of consensus"  
 repeat\_region 21326..21373  
 /note="L1PA6 repeat: matches 6094..6141 of consensus"  
 repeat\_region 22178..22297  
 /note="AluSg/x repeat: matches 1..120 of consensus"  
 repeat\_region 22298..22356  
 /note="Alu repeat: matches 256..308 of consensus"  
 repeat\_region 24152..24454  
 /note="AluY repeat: matches 2..303 of consensus"  
 repeat\_region 25128..25312  
 /note="MER8 repeat: matches 21..214 of consensus"  
 repeat\_region 25330..25517  
 /note="AluJb repeat: matches 108..294 of consensus"  
 repeat\_region 25706..26001  
 /note="AluSx repeat: matches 1..301 of consensus"  
 repeat\_region 26146..26357  
 /note="MER20 repeat: matches 7..218 of consensus"  
 repeat\_region 26399..26685  
 /note="AluSx repeat: matches 2..288 of consensus"  
 repeat\_region 26732..26840  
 /note="L1MC4 repeat: matches 7859..7976 of consensus"  
 repeat\_region 26848..27150  
 /note="AluJb repeat: matches 1..301 of consensus"  
 repeat\_region 27783..28083  
 /note="AluJo repeat: matches 1..298 of consensus"  
 repeat\_region 28086..28394  
 /note="AluSp repeat: matches 1..312 of consensus"  
 repeat\_region 28431..28722  
 /note="AluSg repeat: matches 1..295 of consensus"  
 repeat\_region 29190..29504  
 /note="AluSx repeat: matches 1..311 of consensus"  
 repeat\_region 29685..29734  
 /note="L2 repeat: matches 2648..2697 of consensus"  
 repeat\_region 29759..30061  
 /note="AluJb repeat: matches 1..301 of consensus"  
 repeat\_region 30161..30232  
 /note="AluJ/FRAM repeat: matches 231..302 of consensus"  
 repeat\_region 31184..31483  
 /note="AluSx repeat: matches 1..303 of consensus"  
 repeat\_region 33383..33767  
 /note="AluSc repeat: matches 1..184 of consensus"  
 repeat\_region 33812..34357  
 /note="L2 repeat: matches 1706..2312 of consensus"  
 repeat\_region 35355..36871  
 /note="Cpg island"  
 repeat\_region 35383..35638  
 /note="AluJo repeat: matches 55..312 of consensus"  
 repeat\_region 36676..36817  
 /note="AluJo repeat: matches 6..150 of consensus"  
 repeat\_region 36818..37125  
 /note="AluSg repeat: matches 1..309 of consensus"  
 repeat\_region 37126..37287  
 /note="AluJo repeat: matches 150..303 of consensus"  
 repeat\_region 37322..37614  
 /note="AluSx repeat: matches 1..294 of consensus"  
 repeat\_region 38031..38059  
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Matches 26; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 1 ggcgcgagccattcttcttagcataacggaag 34
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RESULT 15
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DEFINITION Mesorhizobium loti DNA, complete genome, section 16/21.
ACCESSION AP003009 BA000012
VERSION    AP003009.2 GI:14026063
KEYWORDS
SOURCE     Mesorhizobium loti (strain:MAFF303099) DNA.
ORGANISM   Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
            Phyllobacteriaceae; Mesorhizobium.
REFERENCE  1 (sites)
AUTHORS    Kaneko,T., Nakamura,Y., Sato,S., Asamizu,E., Kato,T., Sasamoto,S.,
            Watanabe,A., Idesawa,K., Ishikawa,A., Kawashima,K., Kimura,T.,
            Kishida,Y., Kiyokawa,C., Kohara,M., Matsumoto,M., Matsuno,A.,
            Mochizuki,Y., Nakayama,S., Nakazaki,N., Shimpo,S., Sugimoto,M.,
            Takeuchi,C., Yamada,M. and Tabata,S.
            Complete genome structure of the nitrogen-fixing symbiotic
            bacterium Mesorhizobium loti
            DNA Res. 7 (6), 331-338 (2000)
            21082930
            2 (bases 1 to 339681)
            Kaneko,T.
DIRECT SUBMISSION
Submitted (05-DEC-2000) Takakazu Kaneko, Kazusa DNA Research
Institute, The First Laboratory for Plant Gene Research, Yana
1532-3, Kisarazu, Chiba 292-0812, Japan
(E-mail:kaneko@kazusa.or.jp,
URL:http://www.kazusa.or.jp/rhizobase/,
Tel:81-438-52-3935(ex.2338), Fax:81-438-52-3934)
On May 11, 2001 this sequence version replaced gi:11994984.
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GenCore version 4.5  
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 31, 2002, 17:09:18 ; Search time 2686.26 Seconds  
(without alignments)  
221.075 Million cell updates/sec

Title: US-09-824-567-3

Perfect score: 44

Sequence: 1 ataagaatcgccgcgcacc.....gcaagatatcatggtgggaatc 44

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST.\*

- 1: em\_estba.\*
- 2: em\_esthum.\*
- 3: em\_estin.\*
- 4: em\_estnu.\*
- 5: em\_estov.\*
- 6: em\_estpl.\*
- 7: em\_estro.\*
- 8: em\_hic.\*
- 9: gb\_estl.\*
- 10: gb\_est2.\*
- 11: gb\_hic.\*
- 12: gb\_gss.\*
- 13: em\_gss\_hum.\*
- 14: em\_gss\_inv.\*
- 15: em\_gss\_pln.\*
- 16: em\_gss\_vrt.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
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2	24.6	55.9	505	10 BE442509	BE442509 925021G12
3	24.6	55.9	511	10 BE442509	BE442509 925021G12
4	24.6	55.9	521	10 BE442509	BE442509 925021G12
5	24.6	55.9	613	10 BE442509	BE442509 925021G12
6	24.6	55.9	624	10 BE442509	BE442509 925021G12
7	24.6	55.9	628	10 BE442509	BE442509 925021G12
8	24.6	55.9	638	10 BE442509	BE442509 925021G12
c 9	24.4	55.5	437	9 A1758849	A1758849 ty16b10. x
c 10	24.4	54.5	291	9 B229997	B229997 BE229997
11	23.6	53.6	648	12 AG059299	AG059299 Pan trogl
12	23.4	53.2	648	12 AG059299	AG059299 Pan trogl
13	23.4	53.2	676	12 AG057076	AG057076 Pan trogl
14	23.4	53.2	694	12 AG101006	AG101006 Pan trogl
15	23.2	52.7	452	9 AW547423	AW547423 L002A10
16	23.2	52.7	680	12 AG078608	AG078608 Pan trogl
17	23	52.3	239	9 AV370416	AV370416 AV370416

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c	19	23	52.3	572	10	BM135137	BM135137 WHE0454.D
c	20	23	52.3	582	10	BE417948	BE417948 SCL013.CO
	21	23	52.3	644	12	AG106142	AG106142 Pan trogl
	22	23	52.3	654	12	AG039910	AG039910 Pan trogl
	23	23	52.3	668	12	AG122582	AG122582 Pan trogl
	24	23	52.3	690	12	AG071800	AG071800 Pan trogl
c	25	23	52.3	690	12	AG138371	AG138371 Pan trogl
	26	23	52.3	695	12	AG125018	AG125018 Pan trogl
	27	23	52.3	715	12	AG093362	AG093362 Pan trogl
	28	23	52.3	746	10	BE470112	BE470112 603278786
	29	22.8	51.8	301	10	BG655950	BG655950 IB35B03.Y
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	32	22.8	51.8	648	12	AG033798	AG033798 Pan trogl
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	37	22.6	51.4	720	12	AG052860	AG052860 Pan trogl
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c	42	22.4	50.9	619	10	BE148691	BE148691 602912125
c	43	22.4	50.9	632	11	BC019210	BC019210 Mus muscu
	44	22.4	50.9	647	12	AG036590	AG036590 Pan trogl
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ALIGNMENTS

RESULT 1

BE442509 925021G12.xl C. reinhardtii CC-2290, normalized, Lambda Zap II  
LOCUS Chlamydomonas reinhardtii cDNA, mRNA sequence.  
DEFINITION 493 bp mRNA linear EST 25-JUL-2000  
ACCESSION BE442509.1 GI:9442025  
VERSION EST.  
KEYWORDS Chlamydomonas reinhardtii.  
SOURCE Chlamydomonas reinhardtii  
ORGANISM Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;  
Chlamydomonadaceae; Chlamydomonas.  
REFERENCE 1 (bases 1 to 493)  
AUTHORS Grossman,A., Davies,J., Federspiel,N., Harris,E., Lefebvre,P.,  
McDermott,J.P., Silflow,C., Stern,D. and Surzycki,R.  
TITLE Analyses of the Chlamydomonas reinhardtii genome: A Model,  
Unicellular System for Analyzing Gene Function and Regulation in  
Vascular Plants; project phase 2  
JOURNAL Unpublished (2000)  
COMMENT Contact: Elizabeth H. Harris  
DCMB Box 91000  
Duke University  
Durham, NC 27708-1000, USA  
Tel: 919 613 8164  
Fax: 919 613 8177  
Email: chlamy@duke.edu.

FEATURES

source  
1..493  
/organism="Chlamydomonas reinhardtii"  
/strain="CC-2290 wild type mt- SL D2"  
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/note="Vector: pBluescript II SK-; Site 1: EcoRI; Site 2: XhoI; This library was constructed by John Davies and Jeffrey McDermott. RNA was isolated from strain CC-2290 (Minnesota isolate of C. reinhardtii) grown to mid-log phase in TAP (acetate containing) medium in the light. PolyA mRNA was purified, and cDNA was synthesized and directionally cloned into lambda ZAP II (Stratagene) in

Query Match 55.9%; Score 24.6; DB 10; Length 505;  
Best Local Similarity 76.9%; Pred. No. 48;

RESULT	4
BI720615	521 bp mRNA linear EST 19-SEP-2001
BI720615	1031051A01.x1 C. reinhardtii CC-1690, Stress II (normalized),



Analyses of the *Chlamydomonas reinhardtii* Genome: A Model,  
Unicellular System for Analyzing Gene Function and Regulation in  
Vascular Plants. Project: 1031  
Unpublished (2001)  
Contact: Charles Hauser  
DCMB Box 91000  
Duke University  
Durham, NC 27708-1000  
Tel: 919 613 8159  
Fax: 919 613 8177  
Email: [chauser@duke.edu](mailto:chauser@duke.edu).

cells grown to mid log phase in TAP (NH<sub>4</sub><sup>+</sup> - containing) and shifted to TAP - NO<sub>3</sub><sup>-</sup> (24hrs); H2 production and conditions (0, 12hr, 24hr) see Melis et al., (2000) Plant Phys. 122: 127-135; TAP + H2O2 (1, 12, 24 hr); TAP + sorbitol (1, 2, 6, 24 hr); TAP + Cd (1, 2, 6, 24 hr). PolyA mRNA was purified from each sample, pooled and cDNA synthesized. The cDNA was directionally cloned into lambda Zap II (Stratagene) in the EcoRI (5') and XhoI (3') sites. pBluescript II SK<sup>-</sup> plasmids were excised from the lambda Zap clones by superinfection with EXASS1 (Stratagene) phage. The library was normalized using method 4 described in Bonaldo et al., (1996) Genome

BM001235		624 bp	mRNA	linear	EST 25-OCT-2001
LOCUS					
DEFINITION	x2 C. reinhardtii CC-1690, Stress II (normalized),				
	Lambda zap II Chlamydomonas reinhardtii cDNA, mRNA sequence.				

Chlamydomonas reinhardtii	Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales; Chlamydomonadales; Chlamydomonadales; Chlamydomonas.
1 (bases 1 to 624)	
Grossman,A., Chang,C.-W., Davies,J., Harris,E., Hauser,C., Lefebvre,P., McDermott,J.P., Shrager,J., Siflow,C. and Stern,D.	Analysis of the Chlamydomonas reinhardtii Genome: A Model, Unicellular System for Analyzing Gene Function and Regulation in Vascular Plants. Project: 1031
Unpublished (2001)	
Contact: Charles Hauser	
DCMB Box 91000	
Duke University	
Durham, NC 27708-1000	
Tel: 919 613 8159	
Fax: 919 613 8177	
Email: chauser@duke.edu	

FEATURES  
source

Location/Qualifiers  
1. .624  
/organism="Chlamydomonas reinhardtii"  
/strain="CC-1690 wild type mt+ 21gr"  
/db\_xref="taxon:3055"  
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, Lambda Zap II"  
/note="Vector: pBluescript II SK-; Site\_1: EcoRI; Site\_2:  
XhoI; Stress condition II library, constructed by John  
Davies and Jeffrey McDermott, combines cDNAs from CC-1690  
cells grown to mid-log phase in TAP (NH4+ - containing)  
and shifted to TAP - NO3- (24hrs); H2 production  
conditions (0, 12hr, 24hr) see Melis et al., (2000) Plant  
phys. 122: 127-135; TAP + H2O2 (1, 12, 24 hr); TAP +  
sorbitol (1, 2, 6, 24 hr); TAP + Cd (1, 2, 6, 24 hr).  
PolyA mRNA was purified from each sample, pooled and cDNA  
synthesized. The cDNA was directionally cloned into lambda  
zap II (Stratagene) in the EcoRI (5') and XhoI (3')  
sites. pBluescript II SK- plasmids were excised from the  
lambda ZAP clones by superinfection with ExAssist  
(Stratagene) phage. The library was normalized using  
method 4 described in Bonaldo et al., (1996) Genome  
Research 6: 791-806."  
BASE COUNT 162 a 187 c 139 g 136 t  
ORIGIN

Query Match 55.9%; Score 24.6; DB 10; Length 624;  
Best Local Similarity 76.9%; Pred. No. 51;  
Matches 30; Conservative 0; Mismatches 9; Indels 0; Gaps 0;  
QY 1 ataagaatcgccgcaccatgcgaagatcatcagtg 39  
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Db 445 ATATGAATCGCGCTCAGCTACGCGCAAGTTATTGTGG 483

## RESULT 7

BI723401  
LOCUS BI723401 626 bp mRNA linear EST 19-SEP-2001  
DEFINITION 1031066G11.x3 C. reinhardtii CC-1690, Stress II (normalized),  
Lambda Zap II Chlamydomonas reinhardtii cDNA, mRNA sequence.  
ACCESSION BI723401  
VERSION BI723401.1 GI:15699080  
KEYWORDS EST.  
SOURCE Chlamydomonas reinhardtii.  
ORGANISM Chlamydomonas reinhardtii.  
REFERENCE 1 (bases 1 to 626)  
AUTHORS Grossman,A., Chang,C.-W., Davies,J., Harris,E., Hauser,C., Lefebvre  
P., McDermott,J.P., Shrager,J., Silflow,C. and Stern,D.  
Analyses of the Chlamydomonas reinhardtii Genome: A Model,  
Unicellular System for Analyzing Gene Function and Regulation in  
Vascular Plants. Project: 1031  
JOURNAL Unpublished (2001)  
COMMENT Contact: Charles Hauser  
DCMB Box 91000  
Duke University  
Durham, NC 27708-1000  
Tel: 919 613 8159  
Fax: 919 613 8177  
Email: chauser@duke.edu  
Location/Qualifiers  
1. .626  
/organism="Chlamydomonas reinhardtii"  
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, Lambda Zap II"  
/note="Vector: pBluescript II SK-; Site\_1: EcoRI; Site\_2:  
XhoI; Stress condition II library, constructed by John  
Davies and Jeffrey McDermott, combines cDNAs from CC-1690  
cells grown to mid-log phase in TAP (NH4+ - containing)

FEATURES  
source

Location/Qualifiers  
1. .624  
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, Lambda Zap II"  
/note="Vector: pBluescript II SK-; Site\_1: EcoRI; Site\_2:  
XhoI; Stress condition II library, constructed by John  
Davies and Jeffrey McDermott, combines cDNAs from CC-1690  
cells grown to mid-log phase in TAP (NH4+ - containing)

and shifted to TAP - NO3- (24hrs); H2 production  
conditions (0, 12hr, 24hr) see Melis et al., (2000) Plant  
phys. 122: 127-135; TAP + H2O2 (1, 12, 24 hr); TAP +  
sorbitol (1, 2, 6, 24 hr); TAP + Cd (1, 2, 6, 24 hr).  
PolyA mRNA was purified from each sample, pooled and cDNA  
synthesized. The cDNA was directionally cloned into lambda  
zap II (Stratagene) in the EcoRI (5') and XhoI (3')  
sites. pBluescript II SK- plasmids were excised from the  
lambda ZAP clones by superinfection with ExAssist  
(Stratagene) phage. The library was normalized using  
method 4 described in Bonaldo et al., (1996) Genome  
Research 6: 791-806."  
BASE COUNT 162 a 187 c 140 g 137 t  
ORIGIN

Query Match 55.9%; Score 24.6; DB 10; Length 626;  
Best Local Similarity 76.9%; Pred. No. 51;  
Matches 30; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 1 ataagaatcgccgcaccatgcgaagatcatcagtg 39  
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Db 446 ATATGAATCGCGCTCAGCTACGCGCAAGTTATTGTGG 484

## RESULT 8

BI727233  
LOCUS BI727233 638 bp mRNA linear EST 19-SEP-2001  
DEFINITION 1031090H07.x1 C. reinhardtii CC-1690, Stress II (normalized),  
Lambda Zap II Chlamydomonas reinhardtii cDNA, mRNA sequence.  
ACCESSION BI727233  
VERSION BI727233.1 GI:15702928  
KEYWORDS EST.  
SOURCE Chlamydomonas reinhardtii.  
ORGANISM Chlamydomonas reinhardtii.  
REFERENCE 1 (bases 1 to 638)  
AUTHORS Grossman,A., Chang,C.-W., Davies,J., Harris,E., Hauser,C., Lefebvre  
P., McDermott,J.P., Shrager,J., Silflow,C. and Stern,D.  
Analyses of the Chlamydomonas reinhardtii Genome: A Model,  
Unicellular System for Analyzing Gene Function and Regulation in  
Vascular Plants. Project: 1031  
JOURNAL Unpublished (2001)  
COMMENT Contact: Charles Hauser  
DCMB Box 91000  
Duke University  
Durham, NC 27708-1000  
Tel: 919 613 8159  
Fax: 919 613 8177  
Email: chauser@duke.edu  
Location/Qualifiers  
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/organism="Chlamydomonas reinhardtii"  
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, Lambda Zap II"  
/note="Vector: pBluescript II SK-; Site\_1: EcoRI; Site\_2:  
XhoI; Stress condition II library, constructed by John  
Davies and Jeffrey McDermott, combines cDNAs from CC-1690  
cells grown to mid-log phase in TAP (NH4+ - containing)  
and shifted to TAP - NO3- (24hrs); H2 production  
conditions (0, 12hr, 24hr) see Melis et al., (2000) Plant  
phys. 122: 127-135; TAP + H2O2 (1, 12, 24 hr); TAP +  
sorbitol (1, 2, 6, 24 hr); TAP + Cd (1, 2, 6, 24 hr).  
PolyA mRNA was purified from each sample, pooled and cDNA  
synthesized. The cDNA was directionally cloned into lambda  
zap II (Stratagene) in the EcoRI (5') and XhoI (3')  
sites. pBluescript II SK- plasmids were excised from the  
lambda ZAP clones by superinfection with ExAssist  
(Stratagene) phage. The library was normalized using  
method 4 described in Bonaldo et al., (1996) Genome

Research 6: 791-806."  
BASE COUNT 162 a 187 c 140 g 137 t  
ORIGIN







by long-range high fidelity PCR using Takara's Ex Taq polymerase. Then, the cDNAs were purified by phenol/chloroform and by Centricon 100. The cDNAs were digested with SalI and NotI enzymes. Then, the cDNAs were size selected by Gibco's Size Fractionation Column. The cDNAs were cloned into SalI/NotI site of pSPORT1 plasmid vector. The DH10B E. coli host was transformed with the ligation mixture by chemical method. The library was constructed by Xiaohong Wang."

BASE COUNT 107 a 154 c 127 g 64 t  
ORIGIN

Query Match 52.7%; Score 23.2; DB 9; Length 452;  
Best Local Similarity 77.8%; Pred. No. 1.5e-02;  
Matches 28; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 9 gcggcgccaccatgcgcaagatcatcagtggaatc 44  
| | | | | | | | | | | | | | | | | | | | | |  
Db 411 GGGGGGGGGGACCATGCGCTCAATATCAGGGGGGACTC 446

Search completed: July 31, 2002, 18:59:48  
Job time: 6630 sec

GenCore version 4.5  
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 31, 2002, 17:11:44 ; Search time 84.08 Seconds  
(without alignments)  
128.543 Million cell updates/sec

Title: US-09-824-567-3  
Perfect score: 44  
Sequence: 1 ataagaatgcccgcacc.....gcaagatatcagtggaatc 44

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 383533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued Patents\_NA.\*  
1: /cgn2.6/ptodata/2/ina/5A\_COMB.seq.\*  
2: /cgn2.6/ptodata/2/ina/5B\_COMB.seq.\*  
3: /cgn2.6/ptodata/2/ina/6A\_COMB.seq.\*  
4: /cgn2.6/ptodata/2/ina/6B\_COMB.seq.\*  
5: /cgn2.6/ptodata/2/ina/PCTUS\_COMB.seq.\*  
6: /cgn2.6/ptodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
c 1	22	50.0	414	2	US-08-766-439-20
c 2	22	50.0	414	2	US-08-766-439-21
c 3	22	50.0	1274	2	US-08-766-439-28
c 4	22	50.0	1274	2	US-08-766-439-28
c 5	22	50.0	1327	2	US-08-766-439-29
c 6	22	50.0	1327	2	US-08-766-439-26
c 7	22	50.0	1354	2	US-08-766-439-27
c 8	22	50.0	1354	2	US-08-766-439-24
c 9	20.6	46.8	45	3	US-08-766-439-25
c 10	20.6	46.8	45	4	US-08-974-022-16
c 11	20.6	46.8	45	4	US-08-795-445A-16
c 12	20.6	46.8	45	4	US-08-974-186-16
c 13	20.6	46.8	45	4	US-08-795-445B-16
c 14	20.6	46.8	2412	1	US-08-158-232-9
c 15	20.6	46.8	2412	1	US-08-304-626-9
c 16	20.6	46.8	2412	1	US-08-316-301A-11
c 17	20.6	46.8	2412	1	US-08-611-928-9
c 18	20.6	46.8	2412	3	US-09-173-891-9
c 19	20.6	46.8	2412	4	US-09-076-137-11
c 20	20.6	46.8	2412	5	PCT-US92-03624-11
c 21	20	45.5	35	5	PCT-US96-10905-35
c 22	20	45.5	1491	4	US-09-058-947A-3
c 23	20	45.5	1502	4	US-08-868-373-11
c 24	20	45.5	1807	4	US-09-058-947A-2
c 25	20	45.5	3722	4	US-09-058-947A-1
c 26	19.6	44.5	39	3	US-08-814-052-55
c 27	19.6	44.5	60	5	PCT-US94-08052-5

c 28	19.6	44.5	2249	3	US-08-814-052-19	Sequence 19, Appl
c 29	19.6	44.5	2300	3	US-08-814-052-18	Sequence 18, Appl
c 30	19.4	44.1	75	4	US-09-459-956-20	Sequence 20, Appl
c 31	19.4	44.1	15378	3	US-08-785-420-1	Sequence 1, Appl
c 32	19.2	43.6	1875	1	US-08-453-956-14	Sequence 14, Appl
c 33	19.2	43.6	1875	1	US-08-086-631-14	Sequence 14, Appl
c 34	19.2	43.6	1875	2	US-08-452-930-14	Sequence 14, Appl
c 35	19.2	43.6	1875	5	PCT-US93-08174-14	Sequence 14, Appl
c 36	19.2	43.6	2214	4	US-08-943-731-57	Sequence 57, Appl
c 37	19.2	43.6	2379	4	US-08-797-358B-2	Sequence 2, Appl
c 38	19.2	43.6	18609	4	US-08-943-731-1	Sequence 1, Appl
c 39	19	43.2	44	1	US-08-106-078-7	Sequence 7, Appl
c 40	19	43.2	44	1	US-08-591-492-7	Sequence 24, Appl
c 41	19	43.2	1501	2	US-08-145-658D-24	Sequence 24, Appl
c 42	18.8	42.7	414	2	US-08-766-439-22	Sequence 22, Appl
c 43	18.8	42.7	414	2	US-08-766-439-23	Sequence 23, Appl
c 44	18.8	42.7	560	2	US-08-330-272-3	Sequence 3, Appl
c 45	18.8	42.7	560	5	PCT-US95-13663-3	Sequence 3, Appl

## ALIGNMENTS

RESULT 1  
US-08-766-439-20/c  
; Sequence 20, Application US/08766439  
; Patent No. 5922538  
; GENERAL INFORMATION:  
; APPLICANT: HAZEL, JAMES WILLIAM  
; APPLICANT: JENSEN, MARK ANTON  
; TITLE OF INVENTION: GENETIC MARKERS AND METHODS FOR  
; TITLE OF INVENTION: THE DETECTION OF LISTERIA  
; NUMBER OF SEQUENCES: 110  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: E. I. DU PONT DE NEMOURS AND COMPANY  
; STREET: 1007 MARKET STREET  
; CITY: WILMINGTON  
; STATE: DELAWARE  
; COUNTRY: U.S.A.  
; ZIP: 19898  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: 3.50 INCH DISKETTE  
; OPERATING SYSTEM: MICROSOFT WINDOWS 3.1  
; SOFTWARE: MICROSOFT WORD 2.0C  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/766.439  
; FILING DATE:  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/745,228  
; FILING DATE: NOVEMBER 8, 1996  
; ATTORNEY/AGENT INFORMATION:  
; NAME: FLOYD, LINDA AXAMETHY  
; REGISTRATION NUMBER: 33,692  
; REFERENCE/DOCKET NUMBER: MD-1065-A  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 302-892-8112  
; TELEFAX: 302-773-0164  
; INFORMATION FOR SEQ ID NO: 20:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 414 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; ORIGINAL SOURCE:  
; STRAIN: L MONO - 647 - PREMARKER  
US-08-766-439-20

Query Match 50.0%; Score 22; DB 2; Length 414;

RESULT 3  
US-08-766-439-28/c  
Sequence 28, Application US/08766439  
Patent No. 5922538  
GENERAL INFORMATION:  
APPLICANT: HAZEL, JAMES WILLIAM

ADDRESSER: E. I. DU PONT DE NEMOURS AND COMPANY  
STREET: 1007 MARKET STREET  
CITY: WILMINGTON  
STATE: DELAWARE  
COUNTRY: U.S.A.  
ZIP: 19898  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.50 INCH DISKETTE  
COMPUTER: IBM PC COMPATIBLE



OPERATING SYSTEM: MICROSOFT WINDOWS 3.1  
SOFTWARE: MICROSOFT WORD 2.0C  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/766,439  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION NUMBER: 08/745,228  
FILING DATE: NOVEMBER 8, 1996  
ATTORNEY/AGENT INFORMATION:  
NAME: FLOYD, LINDA AXAMETHY  
REGISTRATION NUMBER: 33,692  
REFERENCE/DOCKET NUMBER: MD-1065-A  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 302-892-8112  
TELEFAX: 302-773-0164  
INFORMATION FOR SEQ ID NO: 29:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1274 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
ANTI-SENSE: YES  
ORIGINAL SOURCE:  
STRAIN: L MONO 3386 D.F.  
US-08-766-439-29

Query Match 50.0%; Score 22; DB 2; Length 1274;  
Best Local Similarity 73.7%; Pred. No. 3.2;  
Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 ataagaatgcgcgcacccatgcgaagatatcagtg 38  
Db 774 AGAAGCATCGCGCGAGATAATGCGCAACTTATTGTG 811

RESULT 5  
US-08-766-439-26/C  
Sequence 26, Application US/08766439  
Patent No. 5922538  
GENERAL INFORMATION:  
APPLICANT: HAZEL, JAMES WILLIAM  
TITLE OF INVENTION: GENETIC MARKERS AND METHODS FOR  
TITLE OF INVENTION: THE DETECTION OF LISTERIA  
TITLE OF INVENTION: MONOCYTOGENES AND LISTERIA SPP.  
NUMBER OF SEQUENCES: 110  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: E. I. DU PONT DE NEMOURS AND COMPANY  
STREET: 1007 MARKET STREET  
CITY: WILMINGTON  
STATE: DELAWARE  
COUNTRY: U.S.A.  
ZIP: 19898  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.50 INCH DISKETTE  
COMPUTER: IBM PC COMPATIBLE  
OPERATING SYSTEM: MICROSOFT WINDOWS 3.1  
SOFTWARE: MICROSOFT WORD 2.0C  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/766,439  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION NUMBER:  
APPLICATION NUMBER: 08/745,228  
FILING DATE: NOVEMBER 8, 1996  
ATTORNEY/AGENT INFORMATION:  
NAME: FLOYD, LINDA AXAMETHY  
REGISTRATION NUMBER: 33,692  
REFERENCE/DOCKET NUMBER: MD-1065-A  
TELECOMMUNICATION INFORMATION:

TELEPHONE: 302-892-8112  
TELEFAX: 302-773-0164  
INFORMATION FOR SEQ ID NO: 26:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1327 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
ORIGINAL SOURCE:  
STRAIN: L MONO 899 D.F.  
US-08-766-439-26

Query Match 50.0%; Score 22; DB 2; Length 1327;  
Best Local Similarity 73.7%; Pred. No. 3.2;  
Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 ataagaatgcgcgcacccatgcgaagatatcagtg 38  
Db 549 AGAAGCATCGCGCGAGATAATGCGCAACTTATTGTG 512

RESULT 6  
US-08-766-439-27  
Sequence 27, Application US/08766439  
Patent No. 5922538  
GENERAL INFORMATION:  
APPLICANT: HAZEL, JAMES WILLIAM  
TITLE OF INVENTION: GENETIC MARKERS AND METHODS FOR  
TITLE OF INVENTION: THE DETECTION OF LISTERIA  
TITLE OF INVENTION: MONOCYTOGENES AND LISTERIA SPP.  
NUMBER OF SEQUENCES: 110  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: E. I. DU PONT DE NEMOURS AND COMPANY  
STREET: 1007 MARKET STREET  
CITY: WILMINGTON  
STATE: DELAWARE  
COUNTRY: U.S.A.  
ZIP: 19898  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.50 INCH DISKETTE  
COMPUTER: IBM PC COMPATIBLE  
OPERATING SYSTEM: MICROSOFT WINDOWS 3.1  
SOFTWARE: MICROSOFT WORD 2.0C  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/766,439  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION NUMBER: 08/745,228  
APPLICATION NUMBER: 08/745,228  
FILING DATE: NOVEMBER 8, 1996  
ATTORNEY/AGENT INFORMATION:  
NAME: FLOYD, LINDA AXAMETHY  
REGISTRATION NUMBER: 33,692  
REFERENCE/DOCKET NUMBER: MD-1065-A  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 302-892-8112  
TELEFAX: 302-773-0164  
INFORMATION FOR SEQ ID NO: 27:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1327 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
ANTI-SENSE: YES  
ORIGINAL SOURCE:  
STRAIN: L MONO 899 D.F.  
US-08-766-439-27

Query Match 50.0%; Score 22; DB 2; Length 1327;  
 Best Local Similarity 73.7%; Pred. No. 3.2;  
 Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ataagaatgcggccaccatgcgaagatatcatg 38  
 Db 779 AGAAGCATGCGCGGAGATATGCGCACTTATTGTG 816

## RESULT 7

US-08-766-439-24/c  
 ; Sequence 24, Application US/08766439  
 ; Patent No. 5922538  
 ; GENERAL INFORMATION:  
 ; APPLICANT: HAZEL, JAMES WILLIAM  
 ; APPLICANT: JENSEN, MARK ANTON  
 ; TITLE OF INVENTION: GENETIC MARKERS AND METHODS FOR  
 ; TITLE OF INVENTION: THE DETECTION OF LISTERIA  
 ; TITLE OF INVENTION: MONOCYTOGENES AND LISTERIA SPP.  
 ; NUMBER OF SEQUENCES: 110  
 ; CORRESPONDENCE ADDRESS:  
 ; ADDRESSEE: E. I. DU PONT DE NEMOURS AND COMPANY  
 ; STREET: 1007 MARKET STREET  
 ; CITY: WILMINGTON  
 ; STATE: DELAWARE  
 ; COUNTRY: U.S.A.  
 ; ZIP: 19898

COMPUTER READABLE FORM:  
 MEDIUM TYPE: 3.50 INCH DISKETTE  
 COMPUTER: IBM PC COMPATIBLE  
 OPERATING SYSTEM: MICROSOFT WINDOWS 3.1  
 SOFTWARE: MICROSOFT WORD 2.0C  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/766,439  
 FILING DATE:

CLASSIFICATION: 435  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 08/745,228  
 FILING DATE: NOVEMBER 8, 1996  
 ATTORNEY/AGENT INFORMATION:  
 NAME: FLOYD, LINDA AXAMETHY  
 REGISTRATION NUMBER: 33,692  
 REFERENCE/DOCKET NUMBER: MD-1065-A  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: 302-892-8112  
 TELEFAX: 302-773-0164  
 INFORMATION FOR SEQ ID NO: 24:

SEQUENCE CHARACTERISTICS:  
 LENGTH: 1354 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA (genomic)  
 ORIGINAL SOURCE:  
 STRAIN: L MONO - 647 - D. FRAG

US-08-766-439-24

Query Match 50.0%; Score 22; DB 2; Length 1354;  
 Best Local Similarity 73.7%; Pred. No. 3.2;  
 Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ataagaatgcggccaccatgcgaagatatcatg 38  
 Db 576 AGAAGCATGCGCGGAGATATGCGCACTTATTGTG 539

## RESULT 8

US-08-766-439-25  
 ; Sequence 25, Application US/08766439  
 ; Patent No. 5922538  
 ; GENERAL INFORMATION:  
 ; APPLICANT: HAZEL, JAMES WILLIAM

APPLICANT: JENSEN, MARK ANTON  
 TITLE OF INVENTION: GENETIC MARKERS AND METHODS FOR  
 TITLE OF INVENTION: THE DETECTION OF LISTERIA  
 TITLE OF INVENTION: MONOCYTOGENES AND LISTERIA SPP.  
 NUMBER OF SEQUENCES: 110  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: E. I. DU PONT DE NEMOURS AND COMPANY  
 STREET: 1007 MARKET STREET  
 CITY: WILMINGTON  
 STATE: DELAWARE  
 COUNTRY: U.S.A.  
 ZIP: 19898

COMPUTER READABLE FORM:  
 MEDIUM TYPE: 3.50 INCH DISKETTE  
 COMPUTER: IBM PC COMPATIBLE  
 OPERATING SYSTEM: MICROSOFT WINDOWS 3.1  
 SOFTWARE: MICROSOFT WORD 2.0C  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/766,439  
 FILING DATE:

CLASSIFICATION: 435  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 08/745,228  
 FILING DATE: NOVEMBER 8, 1996  
 ATTORNEY/AGENT INFORMATION:  
 NAME: FLOYD, LINDA AXAMETHY  
 REGISTRATION NUMBER: 33,692  
 REFERENCE/DOCKET NUMBER: MD-1065-A  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: 302-892-8112  
 TELEFAX: 302-773-0164  
 INFORMATION FOR SEQ ID NO: 25:

SEQUENCE CHARACTERISTICS:  
 LENGTH: 1354 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA (genomic)  
 ANTI-SENSE: YES  
 ORIGINAL SOURCE:  
 STRAIN: L MONO 647 - D.F.  
 US-08-766-439-25

Query Match 50.0%; Score 22; DB 2; Length 1354;  
 Best Local Similarity 73.7%; Pred. No. 3.2;  
 Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ataagaatgcggccaccatgcgaagatatcatg 38  
 Db 779 AGAAGCATGCGCGGAGATATGCGCACTTATTGTG 816

## RESULT 9

US-08-974-022-16  
 ; Sequence 16, Application US/08974022  
 ; Patent No. 6015938  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Boyle, William J.  
 ; APPLICANT: Lacey, David L.  
 ; APPLICANT: Calzone, Frank J.  
 ; APPLICANT: Chang, Ming-Shi  
 ; TITLE OF INVENTION: OSTEOCALCIN  
 ; NUMBER OF SEQUENCES: 53  
 ; CORRESPONDENCE ADDRESS:  
 ; ADDRESSEE: Amgen Inc.  
 ; STREET: 1840 Dehaven Drive  
 ; CITY: Thousand Oaks  
 ; STATE: California  
 ; COUNTRY: USA  
 ; ZIP: 91320-1789  
 ; COMPUTER READABLE FORM:  
 ; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/974,022  
; FILING DATE: 12-DEC-1995  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/577,788  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Winter, Robert B.  
; REFERENCE/DOCKET NUMBER: A-378  
; INFORMATION FOR SEQ ID NO: 16:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 45 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: cDNA  
US-08-974-022-16

Query Match 46.8%; Score 20.6; DB 3; Length 45;  
Best Local Similarity 67.4%; Pred. No. 6.5;  
Matches 29; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 1 ataagaatgcggccgaccatgcgaagatatacagtgggaat 43  
|||||  
Db 1 ATAAGAATGCGCGCGCTAAACTATGAAACAGCCCGAGTGACCAT 43

RESULT 10  
US-08-795-445A-16  
; Sequence 16, Application US/08795445A  
; Patent No. 6284485  
; GENERAL INFORMATION:  
; APPLICANT: Boyle, William J.  
; APPLICANT: Lacey, David L.  
; APPLICANT: Calzone, Frank J.  
; APPLICANT: Chang, Ming-Shi  
; TITLE OF INVENTION: OSTEOPROTEGERIN  
; NUMBER OF SEQUENCES: 53  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Amgen Inc.  
; STREET: 1840 Dehavilland Drive  
; CITY: Thousand Oaks  
; STATE: California  
; COUNTRY: USA  
; ZIP: 91320-1789

; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/795,445A  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/577,788  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Winter, Robert B.  
; REFERENCE/DOCKET NUMBER: A-378  
; INFORMATION FOR SEQ ID NO: 16:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 45 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: cDNA  
US-08-795-445A-16

Query Match 46.8%; Score 20.6; DB 4; Length 45;  
Best Local Similarity 67.4%; Pred. No. 6.5;  
Matches 29; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 1 ataagaatgcggccgaccatgcgaagatatacagtgggaat 43  
|||||  
Db 1 ATAAGAATGCGCGCGCTAAACTATGAAACAGCCCGAGTGACCAT 43

RESULT 11  
US-08-795-447A-16  
; Sequence 16, Application US/08795447A  
; Patent No. 6284728  
; GENERAL INFORMATION:  
; APPLICANT: Boyle, William J.  
; APPLICANT: Lacey, David L.  
; APPLICANT: Calzone, Frank J.  
; APPLICANT: Chang, Ming-Shi  
; TITLE OF INVENTION: Osteoprotegerin  
; NUMBER OF SEQUENCES: 53  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Amgen Inc.  
; STREET: One Amgen Center Drive  
; CITY: Thousand Oaks  
; STATE: California  
; COUNTRY: USA  
; ZIP: 91362-1789  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/795,447A  
; FILING DATE:  
; CLASSIFICATION: 514  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Winter, Robert B.  
; REFERENCE/DOCKET NUMBER: A-378D2  
; INFORMATION FOR SEQ ID NO: 16:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 45 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: cDNA  
US-08-795-447A-16

Query Match 46.8%; Score 20.6; DB 4; Length 45;  
Best Local Similarity 67.4%; Pred. No. 6.5;  
Matches 29; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 1 ataagaatgcggccgaccatgcgaagatatacagtgggaat 43  
|||||  
Db 1 ATAAGAATGCGCGCGCTAAACTATGAAACAGCCCGAGTGACCAT 43

RESULT 12  
US-08-974-186-16  
; Sequence 16, Application US/08974186  
; Patent No. 6284740  
; GENERAL INFORMATION:  
; APPLICANT: Boyle, William J.  
; APPLICANT: Lacey, David L.  
; APPLICANT: Calzone, Frank J.  
; APPLICANT: Chang, Ming-Shi  
; TITLE OF INVENTION: OSTEOPROTEGERIN  
; NUMBER OF SEQUENCES: 53  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Amgen Inc.

STREET: 1840 Dehavilland Drive  
CITY: Thousand Oaks  
STATE: California  
COUNTRY: USA  
ZIP: 91320-1789  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/974,186  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/577,788  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Winter, Robert B.  
REFERENCE/DOCKET NUMBER: A-378  
INFORMATION FOR SEQ ID NO: 16:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 45 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: cDNA  
US-08-974-186-16

Query Match 46.8%; Score 20.6; DB 4; Length 45;  
Best Local Similarity 67.4%; Pred. No. 6.5;  
Matches 29; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 1 atagaatgcgcgcaccatgcgaagatcatcagtggaat 43  
|||||  
Db 1 ATAGAATGCGCGCGCTAAACTATGAACAGCCCGAGTGACCAT 43

RESULT 13  
US-08-795-446B-16  
Sequence 16, Application US/08/795446B  
Patent No. 6288032  
GENERAL INFORMATION:  
APPLICANT: Boyle, William J.  
APPLICANT: Lacey, David L.  
APPLICANT: Calzone, Frank J.  
APPLICANT: Chang, Ming-Shi  
TITLE OF INVENTION: OSTEOPROTEGERIN  
NUMBER OF SEQUENCES: 53  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Amgen Inc.  
STREET: 1840 Dehavilland Drive  
CITY: Thousand Oaks  
STATE: California  
COUNTRY: USA  
ZIP: 91320-1789  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
OPERATING SYSTEM: IBM PC compatible  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/795,446B  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/577,788  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Winter, Robert B.  
REFERENCE/DOCKET NUMBER: A-378  
INFORMATION FOR SEQ ID NO: 16:

SEQUENCE CHARACTERISTICS:  
LENGTH: 45 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: cDNA  
US-08-795-446B-16

Query Match 46.8%; Score 20.6; DB 4; Length 45;  
Best Local Similarity 67.4%; Pred. No. 6.5;  
Matches 29; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 1 atagaatgcgcgcaccatgcgaagatcatcagtggaat 43  
|||||  
Db 1 ATAGAATGCGCGCGCTAAACTATGAACAGCCCGAGTGACCAT 43

RESULT 14  
US-08-158-232-9/c  
Sequence 9, Application US/08158232  
Patent No. 5596071  
GENERAL INFORMATION:  
APPLICANT: Payne, Jewel  
APPLICANT: Kennedy, M. Keith  
APPLICANT: Randall, John Brooks  
APPLICANT: Meier, Henry  
APPLICANT: Dick, Heidi Jane  
APPLICANT: Foncerrada, Luis  
APPLICANT: Schnepf, H. Ernest  
APPLICANT: Schwab, George E.  
APPLICANT: Fu, Jenny  
TITLE OF INVENTION: No. 5596071el Bacillus thuringiensis Toxins Active  
TITLE OF INVENTION: Against Hymenopteran pests  
NUMBER OF SEQUENCES: 51  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: David R. Saliwanchik  
STREET: 2421 N.W. 41st Street, Suite A-1  
CITY: Gainesville  
STATE: FL  
COUNTRY: USA  
ZIP: 32606  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/158,232  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/887,980  
FILING DATE: 22-MAY-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/797,645  
FILING DATE: 25-NOV-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/703,977  
FILING DATE: 22-MAY-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Saliwanchik, David R.  
REGISTRATION NUMBER: 31,794  
REFERENCE/DOCKET NUMBER: M/SCJ104.C1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 904-375-8100  
TELEFAX: 904-372-5800  
INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 2412 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
ORGANISM: Bacillus thuringiensis  
INDIVIDUAL ISOLATE: PS63B  
IMMEDIATE SOURCE:  
CLONE: E. coli NM522(pMYC1642) NRRL B-18961  
US-08-158-232-9

Query Match 46.8%; Score 20.6; DB 1; Length 2412;  
Best Local Similarity 74.3%; Pred. No. 13;  
Matches 26; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 9 gcggccgccaccatgcgcgaagatcagtgaggaaat 43  
||| ||||| || ||||| |||||  
Db 2268 GCGATCGCCACCCACCGAAGATATATTGGGAAT 2234

## RESULT 15

US-08-304-626-9/c  
Sequence 9, Application US/08304626  
Patent No. 5616495  
GENERAL INFORMATION:  
APPLICANT: Payne, Jewel M.  
APPLICANT: Kennedy, M. Keith  
APPLICANT: Randall, John Brooks  
APPLICANT: Meier, Henry  
APPLICANT: Uick, Heidi Jane  
APPLICANT: Foncerrada, Luis  
APPLICANT: Schnepf, Harry E.  
APPLICANT: Schwab, George E.  
TITLE OF INVENTION: No. 5616495el Bacillus thuringiensis Isolates  
TITLE OF INVENTION: Active Against Hymenopteran Pests and Genes Encoding  
TITLE OF INVENTION: Hymenopteran-Active Toxins  
NUMBER OF SEQUENCES: 39  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: David R. Saliwanchik  
STREET: 2421 N.W. 41st Street, Suite A-1  
CITY: Gainesville  
STATE: FL  
COUNTRY: USA  
ZIP: 32606

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/304,626  
FILING DATE:

CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US/07/887,980  
FILING DATE:

ATTORNEY/AGENT INFORMATION:  
NAME: Saliwanchik, David R.  
REGISTRATION NUMBER: 31,794  
REFERENCE/DOCKET NUMBER: M/SCJ 104  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 904-375-8100  
TELEFAX: 904-372-5800

INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 2412 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
ORGANISM: Bacillus thuringiensis  
INDIVIDUAL ISOLATE: PS63B

IMMEDIATE SOURCE:  
CLONE: E. coli NM522(pMYC1642) NRRL B-18961  
US-08-304-626-9

Query Match 46.8%; Score 20.6; DB 1; Length 2412;  
Best Local Similarity 74.3%; Pred. No. 13;  
Matches 26; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 9 gcggccgccaccatgcgcgaagatcagtgaggaaat 43  
||| ||||| || ||||| |||||  
Db 2268 GCGATCGCCACCCACCGAAGATATATTGGGAAT 2234

Search completed: July 31, 2002, 19:32:59  
Job time: 8475 sec

GenCore version 4.5  
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OM nucleic - nucleic search, using sw model

Run on: July 31, 2002, 18:50:09 ; Search time 337.68 Seconds  
(without alignments)  
223.715 Million cell updates/sec

Title: US-09-824-567-3  
Perfect score: 44  
Sequence: 1 ataagaatgcgcgcacc.....gcaagatatcagtggaatc 44

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

c

Database : N\_Geneseq\_032802.\*

- 1: /SID55/gcgdata/geneseq/geneseq-emb1/NA1980.DAT.\*
- 2: /SID55/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.\*
- 3: /SID55/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.\*
- 4: /SID55/gcgdata/geneseq/geneseq-emb1/NA1983.DAT.\*
- 5: /SID55/gcgdata/geneseq/geneseq-emb1/NA1984.DAT.\*
- 6: /SID55/gcgdata/geneseq/geneseq-emb1/NA1985.DAT.\*
- 7: /SID55/gcgdata/geneseq/geneseq-emb1/NA1986.DAT.\*
- 8: /SID55/gcgdata/geneseq/geneseq-emb1/NA1987.DAT.\*
- 9: /SID55/gcgdata/geneseq/geneseq-emb1/NA1988.DAT.\*
- 10: /SID55/gcgdata/geneseq/geneseq-emb1/NA1989.DAT.\*
- 11: /SID55/gcgdata/geneseq/geneseq-emb1/NA1990.DAT.\*
- 12: /SID55/gcgdata/geneseq/geneseq-emb1/NA1991.DAT.\*
- 13: /SID55/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.\*
- 14: /SID55/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.\*
- 15: /SID55/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.\*
- 16: /SID55/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.\*
- 17: /SID55/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.\*
- 18: /SID55/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.\*
- 19: /SID55/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.\*
- 20: /SID55/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.\*
- 21: /SID55/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.\*
- 22: /SID55/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.\*
- 23: /SID55/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.\*
- 24: /SID55/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	44	100.0	44	22	AAD20239
2	27.6	62.7	49	24	AAS18764
3	26.2	59.5	42	24	AAS18778
4	26	59.1	1038602	20	AAS201425
5	25.8	58.6	43	22	AAD09314
6	25.8	58.6	45	22	AAD20895
7	25.4	57.7	51	21	AAS256940
8	25.4	57.7	51	21	AAS256943
9	25	56.8	45	24	AAS18762

10	25	56.8	46	22	AAD09149	Chlamydia pneumoniae
11	24.8	56.4	45	21	AAA75902	PCR primer for DNA
12	24.8	56.4	45	22	AAF83847	C. pneumoniae memb
13	24.8	56.4	45	24	AAS18774	PCR primer #15 use
14	24.8	56.4	48	24	AAS18768	PCR primer #9 used
15	24.6	55.9	43	21	AAS18757	5' PCR primer for
16	24.4	55.5	44	22	AAF83843	C. pneumoniae amin
17	24	54.5	39	21	AAA27123	Chlamydia pneumoniae
18	24	54.5	42	22	AAD20551	C. pneumoniae myos
19	24	54.5	42	22	AAD20879	C. pneumoniae myos
20	24	54.5	42	22	AAF84487	Chlamydia pneumoniae
21	24	54.5	46	24	AAS18770	PCR primer #11 use
22	24	54.5	1235	24	AAS18752	Chlamydia pneumoniae
23	24	54.5	1799	22	AAD20238	Chlamydia pneumoniae
24	24	54.5	1230025	20	AA91990	Nucleotide sequence
25	23.6	53.6	43	21	AAA30923	PCR primer for C.
26	23.6	53.6	46	24	AAS18776	PCR primer #17 use
27	23.6	53.6	2915	23	AAS88477	DNA encoding novel
28	23.4	53.2	43	21	AAD02069	5' primer for ampl
29	23.4	53.2	44	22	AAD03025	Chlamydia pneumoniae
30	23.4	53.2	44	24	AAS18766	PCR primer #7 used
31	23.4	53.2	45	22	AAD20958	C. pneumoniae glut
32	23.4	53.2	45	22	AAF57426	C. pneumoniae lpdA
33	23.2	52.7	43	21	AAD02067	5' primer for ampl
34	23.2	52.7	43	21	AAA75884	PCR primer for DNA
35	23.2	52.7	43	22	AAD20940	Chlamydia pneumoniae
36	23.2	52.7	1638	21	AAC39109	Arabidopsis thalia
37	23	52.3	39	22	AAH46978	Chlamydia general
38	23	52.3	42	21	AAA28409	5' primer for Chla
39	23	52.3	42	21	AAA27019	Chlamydia pneumoniae
40	23	52.3	42	22	AAF31255	Chlamydia pneumoniae
41	23	52.3	42	24	AAS18772	PCR primer #13 use
42	23	52.3	43	21	AAA48840	Chlamydia pneumoniae
43	23	52.3	43	21	AAA28412	5' primer for Chla
44	23	52.3	43	22	AAF89933	PCR primer used to
45	23	52.3	43	24	AAS18760	PCR primer #1 used

ALIGNMENTS

RESULT 1  
AAD20239  
ID AAD20239 standard; DNA; 44 BP.  
XX  
AC AAD20239;  
XX  
DT 15-JAN-2002 (first entry)  
XX  
DE Chlamydia pneumoniae ATP-binding cassette gene amplifying 5'PCR primer.  
XX  
KW ATP-binding cassette; antibiotic; vaccine; infection; therapy; poxvirus;  
KW PCR primer; ss.  
XX  
OS Chlamydia pneumoniae.  
XX  
PN WO200174863-A2.  
XX  
PD 11-OCT-2001.  
XX  
PF 04-APR-2001; 2001WO-CA00455.  
XX  
PR 04-APR-2000; 2000US-194464P.  
XX  
PA (AVET ) AVENTIS PASTEUR LTD.  
XX  
PI Murdin AD, Oomen RP, Wang J, Dunn P;  
XX  
DR WPI; 2001-648549/74.  
XX  
PT Novel Chlamydia ATP-binding cassette and corresponding DNA molecule for  
PT preventing, diagnosing and treating Chlamydia infections in mammals, in  
PT particular humans -

XX Claim 41; Page 53; 89pp; English.  
 XX  
 CC The present invention relates to novel Chlamydia pneumoniae ATP-binding  
 CC cassette protein and its corresponding gene. Sequences of the invention  
 CC are useful for detecting Chlamydia infection by assaying a body fluid  
 CC of a mammal with the components. They are also used as vaccines. ATP  
 CC binding cassette antibodies and vaccines of the invention are useful  
 CC for preventing or treating Chlamydia infection e.g. infection caused  
 CC by C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum in mammals,  
 CC such as humans. The nucleic acid molecules are useful for producing  
 CC ATP-binding cassettes, in the construction of vaccine vectors such  
 CC as poxviruses, which are further useful for preventing and/or treating  
 CC Chlamydia infection and in the construction of attenuated Chlamydia  
 CC strains that can over-express the nucleic acid molecules or express  
 CC it in a non-toxic, mutated form. The present DNA sequence is a 5' PCR  
 CC primer which is used for amplifying Chlamydia pneumoniae ATP-binding  
 CC cassette DNA.  
 XX  
 XX Sequence 44 BP; 14 A; 11 C; 12 G; 7 T; 0 other;

Query Match 100.0%; Score 44; DB 22; Length 44;  
 Best Local Similarity 100.0%; Pred. No. 1e-08;  
 Matches 44; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 ataagaatgccccccaccatgcgcgaagatatcatcagtggaatc 44  
 Db 1 ataagaatgccccccaccatgcgcgaagatatcatcagtggaatc 44

RESULT 2  
 AAS18764  
 ID AAS18764 standard; DNA; 49 BP.  
 XX  
 AC AAS18764;  
 XX  
 DT 26-MAR-2002 (first entry)  
 DE  
 DE PCR primer #5 used to amplify Chlamydia pneumoniae gene.  
 KW ATP binding cassette; secretory locus open reading frame; endopeptidase;  
 KW secretory locus ORF; protease; metalloprotease; CLP protease ATPase;  
 KW CLP protease subunit; transglycolase/transpeptidase; CLP protease;  
 KW thioredoxin; Chlamydia infection; antibacterial; PCR primer; ss.  
 XX  
 OS Chlamydia pneumoniae CWL029.  
 XX  
 PN WO200185972-A2.  
 XX  
 PD 15-NOV-2001.  
 XX  
 PF 08-MAY-2001; 2001WO-CA00653.  
 XX  
 PR 08-MAY-2000; 2000US-202672P.  
 PR 30-MAY-2000; 2000US-207852P.  
 PR 16-JUN-2000; 2000US-211796P.  
 PR 16-JUN-2000; 2000US-211797P.  
 PR 16-JUN-2000; 2000US-211798P.  
 PR 16-JUN-2000; 2000US-211801P.  
 PR 16-JUN-2000; 2000US-212044P.  
 PR 26-SEP-2000; 2000US-235335P.  
 PR 26-SEP-2000; 2000US-235361P.  
 PR 26-SEP-2000; 2000US-235398P.  
 XX  
 PA (AVET ) AVENTIS PASTEUR LTD.  
 XX  
 PI Murdin AD, Oomen RP, Wang J, Dunn P;  
 XX  
 DR WPI; 2002-049447/06.  
 XX  
 PT Vaccine useful for immunising mammals against chlamydia infections,  
 PT comprises vectors having sequences of ATP binding cassette gene,  
 XX

PT secretory locus open reading frame gene of chlamydia -  
 XX  
 PS Example 1; Page 63; 355pp; English.  
 XX  
 CC The present invention relates to the isolation of Chlamydia pneumoniae  
 CC pneumoniae strain CWL029 genes and their encoded proteins. The genes of  
 CC the invention encode an ATP binding cassette gene, a secretory locus  
 CC open reading frame (ORF), an endopeptidase, a protease, a  
 CC metalloprotease, CLP protease ATPase, a CLP protease subunit, a  
 CC transglycolase/transpeptidase, a CLP protease, or thioredoxin. The  
 CC genes of the invention can be used in a vector as a vaccine for the  
 CC prevention and treatment of Chlamydia infections. Also described are  
 CC B- and T-cell epitopes from the proteins of the invention which can be  
 CC used as Chlamydia antigens. AAS18760-AAS18779 represent PCR primers  
 CC used to amplify the C. pneumoniae genes (AAS18750-AAS18759) of the  
 CC invention.  
 XX  
 SQ Sequence 49 BP; 12 A; 14 C; 12 G; 11 T; 0 other;

Query Match 62.7%; Score 27.6; DB 24; Length 49;  
 Best Local Similarity 78.6%; Pred. No. 0.055;  
 Matches 33; Conservative 0; Mismatches 9; Indels 0; Gaps 0;  
 QY 1 ataagaatgccccccaccatgcgcgaagatatcatcagtggaatc 42  
 Db 1 ataagaatgccccccaccatgcgcgaagatatcatcagtggaatc 42

RESULT 3  
 AAS18778  
 ID AAS18778 standard; DNA; 42 BP.  
 XX  
 AC AAS18778;  
 XX  
 DT 26-MAR-2002 (first entry)  
 DE  
 DE PCR primer #19 used to amplify Chlamydia pneumoniae gene.  
 KW ATP binding cassette; secretory locus open reading frame; endopeptidase;  
 KW secretory locus ORF; protease; metalloprotease; CLP protease ATPase;  
 KW CLP protease subunit; transglycolase/transpeptidase; CLP protease;  
 KW thioredoxin; Chlamydia infection; antibacterial; PCR primer; ss.  
 XX  
 OS Chlamydia pneumoniae CWL029.  
 XX  
 PN WO200185972-A2.  
 XX  
 PD 15-NOV-2001.  
 XX  
 PF 08-MAY-2001; 2001WO-CA00653.  
 XX  
 PR 08-MAY-2000; 2000US-202672P.  
 PR 30-MAY-2000; 2000US-207852P.  
 PR 16-JUN-2000; 2000US-211796P.  
 PR 16-JUN-2000; 2000US-211797P.  
 PR 16-JUN-2000; 2000US-211798P.  
 PR 16-JUN-2000; 2000US-211801P.  
 PR 16-JUN-2000; 2000US-212044P.  
 PR 26-SEP-2000; 2000US-235335P.  
 PR 26-SEP-2000; 2000US-235361P.  
 PR 26-SEP-2000; 2000US-235398P.  
 XX  
 PA (AVET ) AVENTIS PASTEUR LTD.  
 XX  
 PI Murdin AD, Oomen RP, Wang J, Dunn P;  
 XX  
 DR WPI; 2002-049447/06.  
 XX  
 PT Vaccine useful for immunising mammals against chlamydia infections,  
 PT comprises vectors having sequences of ATP binding cassette gene,  
 PT secretory locus open reading frame gene of Chlamydia -  
 XX

PS Example 1; Page 68; 355pp; English.

XX The present invention relates to the isolation of Chlamydiaophila  
CC pneumoniae strain CWL029 genes and their encoded proteins. The genes of  
CC the invention encode an ATP binding cassette gene, a secretory locus  
CC open reading frame (ORF), an endopeptidase, a protease, a  
CC metalloprotease, CLP protease Atpase, a CLP protease subunit, a  
CC transglycolase/transpeptidase, a CUPC protease, or thioredoxin. The  
CC genes of the invention can be used in a vector as a vaccine for the  
CC prevention and treatment of Chlamydia infections. Also described are  
CC B- and T-cell epitopes from the proteins of the invention which can be  
CC used as Chlamydia antigens. AAS18760-AAS18779 represent PCR primers  
CC used to amplify the C. pneumoniae genes (AAS18750-AAS18759) of the  
CC invention.

XX Sequence 42 BP; 15 A; 9 C; 10 G; 8 T; 0 other;

Query Match 59.5%; Score 26.2; DB 24; Length 42;  
Best Local Similarity 90.3%; Pred. No. 0.2;  
Matches 28; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 ataagaatgcgcgcaccatgcgaagat 31  
|||||  
Db 1 ataagaatgcgcgcaccatgcgaagat 31

RESULT 4  
AAZ01425/C  
ID AAZ01425 standard; DNA; 1038602 BP.  
AC AAZ01425;  
XX  
DT 07-OCT-1999 (first entry)  
XX  
DE Complete genome sequence of Chlamydia trachomatis.  
XX

Vaccine; eye disease; conventional trachoma; nonendemic trachoma;  
KW paratrachoma; inclusion conjunctivitis; genital disease; perinephritis;  
KW nongonococcal urethritis; epididymitis; cervicitis; salpingitis;  
KW Bartholinitis; pneumonia; venereal lymphogranulomatosis; ss.  
XX Chlamydia trachomatis.

XX W0928475-A2.  
XX  
XX 10-JUN-1999.

XX 27-NOV-1998; 98WO-IB01939.  
XX  
XX 04-NOV-1998; 98US-0107077.  
XX 28-NOV-1997; 97FR-0015041.  
XX 17-DEC-1997; 97FR-0016034.

XX (GENT ) GENSET.

XX Griffais R;

XX WPI; 1999-371125/31.

XX Genome sequence of Chlamydia trachomatis  
XX  
XX Claim 1; Page 373-656; 1755pp; English.

XX The present sequence represents the complete genome of Chlamydia  
CC trachomatis. Open reading frames (ORFs) of the genome encode  
CC polypeptides AAY36754-Y37949. The polypeptides can be used as vaccines  
CC against Chlamydia trachomatis. Antisense and ribozyme sequences can also  
CC be used to control growth of the microorganism. Chlamydia trachomatis is  
CC responsible for a large number of diseases, e.g. eye diseases such as  
CC conventional trachoma, nonendemic trachoma, paratrachoma, and inclusion  
CC conjunctivitis; genital diseases such as nongonococcal urethritis,  
CC epididymitis, cervicitis, salpingitis, perinephritis, Bartholinitis;

CC pneumonia in breast feeding infants; and venereal  
CC lymphogranulomatosis. The polypeptides of the invention may be of use in  
CC treating these diseases.

XX Sequence 1038602 BP; 304265 A; 214645 C; 214259 G; 305001 T; 432 other;

Query Match 59.1%; Score 26; DB 20; Length 1038602;  
Best Local Similarity 100.0%; Pred. No. 2;  
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 19 ccacgcgaagatcatcagtggaatc 44  
|||||  
Db 252099 CCATCCGCAAGATATCATGCGGATC 252074

RESULT 5  
AAD09314  
ID AAD09314 standard; DNA; 43 BP.  
XX  
AC AAD09314;  
XX  
DT 10-SEP-2001 (first entry)  
XX  
DE Chlamydia pneumoniae outer membrane protein gene amplifying 5'PCR primer.  
XX  
KW Outer membrane protein; therapy; Chlamydia infection;  
KW antibiotic; vaccine; PCR primer; ss.  
XX Chlamydia pneumoniae.

XX W0200146225-A2.  
XX  
XX 28-JUN-2001.

XX 20-DEC-2000; 2000WO-CA01535.  
XX  
XX 22-DEC-1999; 99US-0171539.

XX (AVET ) AVENTIS PASTEUR LTD.  
XX  
XX Murdin AD, Oomen RP, Wang J, Dunn P;

XX WPI; 2001-418020/44.  
XX  
XX Chlamydia outer membrane protein and corresponding DNA molecules for  
XX preventing, diagnosing and treating Chlamydia infection in mammals,  
XX such as humans -

XX Claim 32; Page 52; 74pp; English.  
XX  
XX The present sequence is a PCR primer which is used to amplify the  
XX Chlamydia pneumoniae outer membrane protein gene. The outer membrane  
XX protein is useful for preventing, treating and detecting Chlamydia  
XX infection in humans. The outer membrane protein DNA is useful for  
XX producing the encoded polypeptide and in the construction of attenuated  
XX Chlamydia strains that can over express the polynucleotide or express  
XX it in a non-toxic, mutated form. It is also used as vaccine. The probes  
XX for outer membrane protein are useful in diagnostic tests as capture or  
XX detection probes and the primers are useful in diagnostic methods  
XX involving PCR. The antibody against outer membrane protein is useful for  
XX purifying the outer membrane protein.

XX Sequence 43 BP; 16 A; 9 C; 6 G; 12 T; 0 other;

Query Match 58.6%; Score 25.8; DB 22; Length 43;  
Best Local Similarity 81.1%; Pred. No. 0.29;  
Matches 30; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

Qy 1 ataagaatgcgcgcaccatgcgaagatcagt 37  
|||||  
Db 1 ataagaatgcgcgcaccatgaaaaattattt 37



```

RESULT 6
AAD20895
ID AAD20895 standard; DNA; 45 BP.
XX AC
XX AAD20895;
XX DT
XX 15-JAN-2002 (first entry)
XX DE
XX C. pneumoniae myosin gene amplifying 5' PCR primer alternative version.
XX KW Transmembrane protein; antibacterial; vaccine; gene therapy;
XX KW immunisation; PCR primer; ss.
XX OS Chlamydia pneumoniae.
XX PA WO200175114-A2.
XX PN 11-OCT-2001.
XX PD
XX PF 04-APR-2001; 2001WO-CR00462.
XX PR
XX 04-APR-2000; 2000US-194477P.
XX PA (AVET ) AVENTIS PASTEUR LTD.
XX PI Murdin AD, Oomen RP, Wang J, Dunn P;
XX XX WPI; 2001-648559/74.
XX XX
XX PT Novel polypeptides from Chlamydia pneumoniae and genes encoding the
XX PT polypeptide, useful for immunization of host e.g. human against disease
XX PT caused by infection by a strain of Chlamydia
XX PS Claim 41; Page 89; 90pp; English.
XX XX
XX CC The invention relates to a transmembrane polypeptide from Chlamydia,
XX CC preferably C. pneumoniae. Transmembrane protein and its gene are useful
XX CC as vaccines and for preventing or treating Chlamydia infection.
XX CC Transmembrane protein, its gene and antibody are useful for detecting
XX CC Chlamydia infection, by assaying a body fluid of a mammal to be tested
XX CC The probes are used in diagnostic tests as capture or detection probes
XX CC and in hybridisation techniques, and primers are useful in amplification
XX CC techniques for use in diagnostic methods. Transmembrane protein is useful
XX CC for detecting the presence of anti-Chlamydia antibodies in blood sample.
XX CC The present sequence is an alternative version of a PCR primer used for
XX CC amplifying C. pneumoniae myosin heavy chain homologue gene used in the
XX CC exemplification of the invention.
XX CC Note: This sequence is stated as being the same as SEQ ID NO: 3 shown
XX CC in page 53 (AAD20879) of the specification. However, the sequences differ
XX CC at several positions.
XX SQ Sequence 45 BP; 12 A; 14 C; 8 G; 11 T; 0 other;

Query Match 58.6%; Score 25.8; DB 22; Length 45;
Best Local Similarity 93.1%; Pred. No. 0.3;
Matches 27; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ataagaatgcggcgccaccatgcgcaag 29
Db 1 ataagaatgcggcgccaccatgcgcaag 29
|||||
RESULT 7
AAZ56940
ID AAZ56940 standard; DNA; 51 BP.
XX AC
XX AAZ56940;
XX XX
XX 08-MAY-2000 (first entry)
XX DT
XX PF
XX 27-JUL-1999; 99WO-IB01333.
XX XX

C. pneumoniae mip gene amplifying 5' primer.
Chlamydia pneumoniae; outer membrane protein; mip; CPN100501;
Chlamydial infection; PCR primer; ss.
Chlamydia pneumoniae.
WO200006741-A1.
10-FEB-2000.
27-JUL-1999; 99WO-IB01330.
27-JUL-1998; 98US-0094192.
01-MAR-1999; 98US-0122044.
26-JUL-1999; 98US-0361440.
(CONN-) CONNAUGHT LAB LTD.
Murdin AD, Oomen RP, Dunn PL;
WPI; 2000-195302/17.
Novel polynucleotides and Chlamydia pneumoniae outer membrane protein
encoded by them for use as vaccines in treating and diagnosing
chlamydial infections
Example 1; Page 34; 55pp; English.
The invention provides an isolated polynucleotide encoding Chlamydia
pneumoniae outer membrane protein (mip or CPN100501). The mip protein
can be expressed by standard recombinant methodology. The mip gene is
used for detecting Chlamydia by hybridizing or amplifying the sample
with the mip gene specific probe. A vaccine vector or a pharmaceutical
composition comprising the mip gene are used for inducing an immune
response in a mammal to prevent/treat chlamydial infections particularly
infections caused by C. pneumoniae. The present sequence represents the
a PCR primer amplifying the C. pneumoniae mip gene.
Sequence 51 BP; 20 A; 9 C; 9 G; 13 T; 0 other;

Query Match 57.7%; Score 25.4; DB 21; Length 51;
Best Local Similarity 82.9%; Pred. No. 0.44;
Matches 29; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 ataagaatgcggcgccaccatgcgcaagatcatca 35
Db 1 ataagaatgcggcgccaccatgcgcaagatcatca 35
|||||
RESULT 8
AAZ56943
ID AAZ56943 standard; DNA; 51 BP.
XX AC
XX AAZ56943;
XX DT
XX 08-MAY-2000 (first entry)
XX DE
XX C. pneumoniae omp gene amplifying 5' primer.
XX KW Chlamydia pneumoniae antigen; omp; CPN100314; antibacterial; PCR primer;
XX KW vaccination; Chlamydia infection; community acquired pneumonia;
XX KW upper respiratory tract infection; bronchitis; sinusitis; ss.
XX OS Chlamydia pneumoniae.
XX PN WO200006743-A2.
XX PD
XX 10-FEB-2000.
XX PF
XX 27-JUL-1999; 99WO-IB01333.
XX XX

```



PA (AVET ) AVENTIS PASTEUR LTD.  
 PI Murdin AD, Oomen RP, Wang J, Dunn P;  
 XX WPI; 2001-418021/44.  
 DR Chlamydia polypeptides, designated membrane ATPase and corresponding  
 PT DNA molecules for preventing, diagnosing and treating Chlamydia  
 PT infection in mammals, including humans -  
 XX Claim 32; Page 52; 80pp; English.  
 XX The present sequence is a 5' PCR primer used to amplify  
 CC Chlamydia pneumoniae membrane ATPase gene. Membrane ATPase is used  
 CC as vaccine. Membrane ATPase is useful for detecting, preventing and  
 CC treating Chlamydia infection such as pneumoniae, bronchitis, sinusitis,  
 CC acute respiratory disease, cough, fever, abnormal chest sounds, lower  
 CC respiratory tract infection, atherosclerosis and asthma, in mammals,  
 CC in particular humans. Membrane ATPase is useful in the construction of  
 CC attenuated Chlamydia strains that can over express the polynucleotide or  
 CC express it in a non-toxic, mutated form.  
 XX Sequence 46 BP; 21 A; 13 C; 6 G; 6 T; 0 other;  
 SQ

Query Match 56.8%; Score 25; DB 22; Length 46;  
 Best Local Similarity 84.8%; Pred. No. 0.63;  
 Matches 28; Conservative 0; Mismatches 5; Indels 0; Gaps 0;  
 QY 1 ataagaatgcgcgcaccatgcgaagatat 33  
 DB 1 ataagaatgcgcgcaccatgcgaacaaatct 33

RESULT 11  
 AAA75902  
 ID AAA75902 standard; DNA; 45 BP.  
 XX AAA75902;  
 AC  
 XX 22-JAN-2001 (first entry)  
 DT  
 XX PCR primer for DNA encoding a 60 kda cysteine-rich membrane protein.  
 DE  
 XX Cysteine-rich membrane protein; Chlamydia infection; bronchitis;  
 KW community acquired pneumonia; upper respiratory tract infection; vaccine;  
 KW sinusitis; PCR primer; ss.  
 XX Chlamydia pneumoniae.  
 OS  
 XX WO200055326-A1.  
 PN  
 XX 21-SEP-2000.  
 PD  
 XX 09-MAR-2000; 2000WO-CA00240.  
 PF  
 XX 12-MAR-1999; 99US-0123966.  
 PR  
 XX (AVET ) AVENTIS PASTEUR LTD.  
 PA  
 XX Murdin AD, Oomen RP, Wang J, Dunn P;  
 PI WPI; 2000-618918/59.  
 XX New polynucleotides encoding a 60kda cysteine-rich membrane protein  
 PT from Chlamydia, useful as a vaccine for preventing and treating  
 PT Chlamydia infection in mammals -  
 XX Example 1; Page 48; 77pp; English.  
 XX PCR primers AAA75902-03 were used to amplify DNA encoding a Chlamydia  
 CC 60 kda cysteine-rich membrane protein. The membrane-rich polynucleotide  
 CC and polypeptide are useful for preventing or treating Chlamydia

CC infection, such as community pneumonia, upper respiratory tract  
 CC infections, bronchitis and sinusitis. They are also useful for  
 CC diagnosing Chlamydia infection by assaying a body fluid of a mammal.  
 CC The polypeptide is useful for vaccine production.  
 XX Sequence 45 BP; 15 A; 13 C; 10 G; 7 T; 0 other;  
 SQ

Query Match 56.4%; Score 24.8; DB 21; Length 45;  
 Best Local Similarity 92.9%; Pred. No. 0.76;  
 Matches 26; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 ataagaatgcgcgcaccatgcgcaa 28  
 DB 1 ataagaatgcgcgcaccatgcgcaa 28

RESULT 12  
 AAF83847  
 ID AAF83847 standard; DNA; 45 BP.  
 XX AAF83847;  
 AC  
 XX 06-AUG-2001 (first entry)  
 DT  
 XX C. pneumoniae membrane ATPase DNA amplifying 5' primer.  
 DE  
 XX Membrane adenosine triphosphatase; ATP; membrane ATPase; Chlamydia;  
 KW infection; antibiotic; vaccine; PCR primer; ss.  
 KW Chlamydia pneumoniae.  
 OS  
 XX WO200136455-A2.  
 PN  
 XX 25-MAY-2001.  
 PD  
 XX 10-NOV-2000; 2000WO-CA01344.  
 PF  
 XX 12-NOV-1999; 99US-0164823.  
 PR  
 XX (AVET ) AVENTIS PASTEUR LTD.  
 PA  
 XX Murdin AD, Oomen RP, Wang J, Dunn P;  
 PI WPI; 2001-355610/37.  
 DR  
 XX Novel Chlamydia pneumoniae membrane ATPase polypeptide useful for  
 PT preventing, treating or detecting Chlamydia infection -  
 PT Claim 32; Page 52; 83pp; English.  
 PS  
 XX Sequences AAF83847-848 represent PCR primers for amplifying the genomic  
 CC DNA encoding a Chlamydia pneumoniae membrane adenosine triphosphate  
 CC (ATPase) polypeptide. The membrane ATPase polynucleotides and polypeptide  
 CC and fragments are useful for preventing or treating Chlamydia infection  
 CC and for detecting Chlamydia by assaying a body fluid. The polynucleotides  
 CC are useful for producing recombinant membrane ATPase, in the construction  
 CC of vaccine vectors for preventing and/or treating Chlamydia infection,  
 CC as a vaccine agent, and in the construction of attenuated Chlamydia  
 CC strains that can over-express the polypeptide, or express it in a  
 CC non-toxic, mutated form.  
 XX Sequence 45 BP; 16 A; 11 C; 10 G; 8 T; 0 other;  
 SQ

Query Match 56.4%; Score 24.8; DB 22; Length 45;  
 Best Local Similarity 80.6%; Pred. No. 0.76;  
 Matches 29; Conservative 0; Mismatches 7; Indels 0; Gaps 0;  
 QY 1 ataagaatgcgcgcaccatgcgaagatatcag 36  
 DB 1 ataagaatgcgcgcaccatggttaacagtttcag 36

## RESULT 13

AAS18774  
ID AAS18774 standard; DNA; 45 BP.

XX AAS18774;  
XX

XX 26-MAR-2002 (first entry)  
XX

XX PCR primer #15 used to amplify Chlamydomophila pneumoniae gene.  
XX

XX ATP binding cassette; secretory locus open reading frame; endopeptidase;  
XX secretory locus ORF; protease; metalloprotease; CLP protease ATPase;  
KW CLP protease subunit; transglycolase/transpeptidase; CLPc protease;  
KW thiorodoxin; Chlamydia infection; antibacterial; PCR primer; ss.  
XX

OS Chlamydomophila pneumoniae CWL029.  
XX

XX WO200185972-A2.  
XX

XX 15-NOV-2001.  
XX

XX 08-MAY-2001; 2001WO-CA00653.  
XX

XX 08-MAY-2000; 2000US-202672P.  
XX

XX 30-MAY-2000; 2000US-207852P.  
XX

XX 16-JUN-2000; 2000US-211796P.  
XX

XX 16-JUN-2000; 2000US-211797P.  
XX

XX 16-JUN-2000; 2000US-211798P.  
XX

XX 16-JUN-2000; 2000US-212044P.  
XX

XX 26-SEP-2000; 2000US-235335P.  
XX

XX 26-SEP-2000; 2000US-235361P.  
XX

XX 26-SEP-2000; 2000US-235398P.  
XX

XX (AVET ) AVENTIS PASTEUR LTD.  
XX

XX Murdin AD, Oomen RP, Wang J, Dunn P;  
XX

XX WPI; 2002-049447/06.  
XX

XX Vaccine useful for immunising mammals against chlamydia infections,  
XX comprises vectors having sequences of ATP binding cassette gene,  
XX secretory locus open reading frame gene of chlamydia -  
XX

XX Example 1; Page 66; 355pp; English.  
XX

XX The present invention relates to the isolation of Chlamydomophila  
XX pneumoniae strain CWL029 genes and their encoded proteins. The genes of  
XX the invention encode an ATP binding cassette gene, a secretory locus  
XX open reading frame (ORF), an endopeptidase, a protease, a  
XX metalloprotease, CLP protease ATPase, a CLP protease subunit, a  
XX transglycolase/transpeptidase, a CLPc protease, or thiorodoxin. The  
XX genes of the invention can be used in a vector as a vaccine for the  
XX prevention and treatment of Chlamydia infections. Also described are  
XX B- and T-cell epitopes from the proteins of the invention which can be  
XX used as Chlamydia antigens. AAS18760-AAS18779 represent PCR primers  
XX used to amplify the C. pneumoniae genes (AAS18750-AAS18759) of the  
XX invention.  
XX

XX Sequence 45 BP; 13 A; 14 C; 10 G; 8 T; 0 other;  
SQ

Query Match 56.4%; Score 24.8; DB 24; Length 45;  
Best Local Similarity 92.9%; Pred. No. 0.76;  
Matches 26; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ataagaatgcggccgaccatgcgcaa 28  
|||||

Db 1 ataagaatgcggccgaccatgcgcta 28  
|||||

## RESULT 14

AAS187587  
ID AAS187587 standard; DNA; 43 BP.

## AAS18768

XX AAS18768 standard; DNA; 48 BP.

XX AAS18768;  
XX

XX 26-MAR-2002 (first entry)  
XX

XX PCR primer #9 used to amplify Chlamydomophila pneumoniae gene.  
XX

XX ATP binding cassette; secretory locus open reading frame; endopeptidase;  
XX secretory locus ORF; protease; metalloprotease; CLP protease ATPase;  
KW CLP protease subunit; transglycolase/transpeptidase; CLPc protease;  
KW thiorodoxin; Chlamydia infection; antibacterial; PCR primer; ss.  
XX

OS Chlamydomophila pneumoniae CWL029.  
XX

XX WO200185972-A2.  
XX

XX 15-NOV-2001.  
XX

XX 08-MAY-2001; 2001WO-CA00653.  
XX

XX 08-MAY-2000; 2000US-202672P.  
XX

XX 30-MAY-2000; 2000US-207852P.  
XX

XX 16-JUN-2000; 2000US-211796P.  
XX

XX 16-JUN-2000; 2000US-211797P.  
XX

XX 16-JUN-2000; 2000US-211798P.  
XX

XX 16-JUN-2000; 2000US-211801P.  
XX

XX 26-SEP-2000; 2000US-235335P.  
XX

XX 26-SEP-2000; 2000US-235361P.  
XX

XX 26-SEP-2000; 2000US-235398P.  
XX

XX (AVET ) AVENTIS PASTEUR LTD.  
XX

XX Murdin AD, Oomen RP, Wang J, Dunn P;  
XX

XX WPI; 2002-049447/06.  
XX

XX Vaccine useful for immunising mammals against chlamydia infections,  
XX comprises vectors having sequences of ATP binding cassette gene,  
XX secretory locus open reading frame gene of chlamydia -  
XX

XX Example 1; Page 64; 355pp; English.  
XX

XX The present invention relates to the isolation of Chlamydomophila  
XX pneumoniae strain CWL029 genes and their encoded proteins. The genes of  
XX the invention encode an ATP binding cassette gene, a secretory locus  
XX open reading frame (ORF), an endopeptidase, a protease, a  
XX metalloprotease, CLP protease ATPase, a CLP protease subunit, a  
XX transglycolase/transpeptidase, a CLPc protease, or thiorodoxin. The  
XX genes of the invention can be used in a vector as a vaccine for the  
XX prevention and treatment of Chlamydia infections. Also described are  
XX B- and T-cell epitopes from the proteins of the invention which can be  
XX used as Chlamydia antigens. AAS18760-AAS18779 represent PCR primers  
XX used to amplify the C. pneumoniae genes (AAS18750-AAS18759) of the  
XX invention.  
XX

XX Sequence 48 BP; 17 A; 11 C; 8 G; 12 T; 0 other;  
SQ

Query Match 56.4%; Score 24.8; DB 24; Length 48;  
Best Local Similarity 92.9%; Pred. No. 0.77;  
Matches 26; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ataagaatgcggccgaccatgcgcaa 28  
|||||

Db 1 ataagaatgcggccgaccatgcgaaa 28  
|||||

XX AAZ61587;  
XX 19-JUN-2000 (first entry)  
XX  
XX 5' PCR primer for POMP91A gene of Chlamydia pneumoniae.  
XX POMP91A; Chlamydia pneumoniae strain CMI; Chlamydia infection;  
XX vaccine; immune response; PCR primer; ss.  
XX Chlamydia pneumoniae.  
XX  
XX WO200011180-A1.  
XX  
XX 02-MAR-2000.  
XX  
XX 19-AUG-1999; 99WO-CA00765.  
XX  
XX 20-AUG-1998; 98US-0097198.  
XX  
XX (CONN-) CONNAUGHT LAB LTD.  
XX  
XX Murdin AD, Dunn PL, Oomen RP;  
XX  
XX WPI; 2000-224700/19.  
XX  
XX New nucleic acid encoding POMP91A protein from a strain of Chlamydia  
XX useful for preventing, treating and diagnosing Chlamydia infection -  
XX  
XX Example; Page 38; 98pp; English.

XX PCR primers AAZ61587-88 were used to amplify DNA encoding a polypeptide  
XX of POMP91A from Chlamydia pneumoniae strain CMI genomic DNA. The  
XX polynucleotides or polypeptides are used to prevent, treat and  
XX diagnose Chlamydia infection. Vaccine vectors containing POMP91A  
XX polynucleotides are used to induce an immune response against  
XX Chlamydia. Antibodies against POMP91A can be used to diagnose the  
XX presence of Chlamydia in a biological sample.  
XX  
XX Sequence 43 BP; 11 A; 10 C; 14 G; 8 T; 0 other;

Query Match 55.9%; Score 24.6; DB 21; Length 43;  
Best Local Similarity 87.1%; Pred. No. 0.91;  
Matches 27; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 1 ataagaatcgccgccaccatgccaagat 31  
Db 1 ataagaatcgccgccaccatgccaagat 31  
|||||

Search completed: July 31, 2002, 20:58:36  
Job time: 7707 sec

OM of: US-09-824-567-2 to: EST:\* out\_format : pfs

Date: Jul 26, 2002 5:12 AM

About: Results were produced by the GenCore software, version 4.5,  
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Command line parameters:

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-MINMATCH=0.100 -LOOPL=0.000 -LOOPEXT=0.000 -QGAPO=4.500
-OGAPEXT=0.050 -XGAPO=10.000 -XGAPEXT=0.500 -FGAPO=6.000
-OGAPEXT=7.000 -YGAPO=10.000 -YGAPEXT=0.500 -DELOP=6.000
-DELEXT=7.000 -START=1 -MATRIX=blotsum62 -TRANS=human40.cdi
-LIST=45 -DOALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0
-ALIGN=15 -MODE=LOCAL -OUTFMT=pfs -NORM=ext -HEAPSIZ=500
-MINLEN=0 -MAXLEN=200000000 -USER=US09824567 @CNC1_1.5096
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-NO_XUPXY -WAIT -THREADS=1
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Search information block:

Query: US-09-824-567-2

Query length: 532

Database: EST\*

Database sequences: 13736207

Database length: -1841457050

Search time (sec): 1769.610000

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gb_gss:AO990639	+	148.50	266.40	1.4e-05	538	AO990639 RfC01442 Photornabidus
gb_gss:BA370951	-	137.00	243.09	0.0003	593	BA370951 AG-ND-119H23 TF ND-TAM
gb_gss:BO7758	-	124.50	218.87	0.0061	601	BO7758 8219P101E0112019577 Rhod
gb_gss:AO990866	-	122.50	214.78	0.0102	613	AO990866 GSSBrl1690 Photornabidus
gb_gss:AC302752	+	118.00	211.22	0.0162	402	AC302752 GSSBrl1690 Brucella ab
gb_gss:AO012177	+	114.50	197.56	0.0932	713	AO012177 2711C073112697 Cosmid
gb_est1:AI503668	+	109.00	194.26	0.1421	390	AI503668 VK75e04.X1 Knowles Sol
gb_gss:AO579158	+	108.50	187.24	0.3499	644	AO579158 nbxb0096M13f CUGI Rice
gb_gss:BO7745	-	104.00	181.68	0.7133	498	BO7745 8025R101A07082895T3 Rhod
gb_gss:TA38006P	+	103.50	180.22	0.8603	519	TA38006P T. brucei sheared gene
gb_gss:BA398784	-	103.50	179.43	0.9519	554	BA398784 AG-ND-150B19 TR ND-TAM
gb_gss:TA358F03P	-	103.50	179.35	0.9626	558	TA358F03P T. brucei sheared gene
gb_gss:AC302822	+	102.50	176.89	1.32	391	AC302822 GSSBrl1690 Brucella ab
gb_gss:BF574618	-	102.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
gb_est2:BI180815	-	102.00	169.88	3.24	962	BI180815 3121C3 LRH (Lin Rhoda
gb_est2:BI180755	+	101.50	179.95	0.8908	386	BI180755 LRH21C3 LRH (Lin Rhoda
gb_est2:BE73898	+	101.50	179.89	0.8979	388	BE73898 602133277F1 NIH_MGC_81
gb_est2:BF574618	-	101.50	176.89	1.32	583	BF574618 AG-ND-17AG16 TR ND-TAM
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GenCore version 4.5  
Copyright (c) 1993 - 2000 Compugen Ltd.

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Listing first 45 summaries

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32: em\_hgt\_other.\*

33: em\_hgt\_inv.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

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3	26.2	59.5	42	6	AX300427	Sequence
4	26	59.1	11944	1	AE001293	Chlamydia
5	25.8	58.6	43	6	AX179663	Sequence
6	25.8	58.6	45	6	AX278234	Sequence
7	25	56.8	45	6	AX300411	Sequence
8	25	56.8	46	6	AX179705	Sequence
9	24.8	56.4	45	6	AX146974	Sequence
10	24.8	56.4	45	6	AX300423	Sequence
11	24.8	56.4	48	6	AX300417	Sequence
12	24.4	55.5	44	6	AX147161	Sequence
13	24.4	55.5	12173	1	AE002315	Chlamydia
14	24	54.5	42	6	AX100501	Sequence
15	24	54.5	42	6	AX268467	Sequence
16	24	54.5	46	6	AX300419	Sequence
17	24	54.5	1235	6	AX300393	Sequence
18	24	54.5	1599	6	AX349501	Sequence
19	24	54.5	1799	6	AX268341	Sequence
20	24	54.5	11648	1	AE001606	Chlamydia
21	24	54.5	11764	1	AE002216	Chlamydia
22	24	54.5	299650	1	AP002545	Chlamydia
23	23.6	53.6	46	6	AX300425	Sequence
24	23.4	53.2	43	6	AX045141	Sequence
25	23.4	53.2	44	6	AX100530	Sequence
26	23.4	53.2	44	6	AX300415	Sequence
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ACCESSION	AX268343	AX268343	Sequence 3 from Patent WO0174863.	44 bp	DNA	linear	PAT 29-OCT-2001
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Result No.

Score

Query Match

Length

ID

Description

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 VERSION AX300413.1 GI:17381804  
 KEYWORDS synthetic construct.  
 SOURCE synthetic construct.  
 ORGANISM artificial sequence.  
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 AUTHORS Mardin, A.D., Omen, R.P., Wang, J. and Dunn, P.  
 TITLE Chlamydia antigens and corresponding dna fragments and uses thereof  
 JOURNAL Patent: WO 0185972-A 25 15-NOV-2001;  
 Aventis Pasteur Limited (CA)  
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 VERSION AX300427.1 GI:17381818  
 KEYWORDS synthetic construct.  
 SOURCE synthetic construct.  
 ORGANISM artificial sequence.  
 REFERENCE 1 (sites)  
 AUTHORS Mardin, A.D., Omen, R.P., Wang, J. and Dunn, P.  
 TITLE Chlamydia antigens and corresponding dna fragments and uses thereof  
 JOURNAL Patent: WO 0185972-A 39 15-NOV-2001;  
 Aventis Pasteur Limited (CA)  
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 VERSION AE001293.1 GI:3328597  
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 SOURCE Chlamydia trachomatis.  
 ORGANISM Chlamydia trachomatis.  
 Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.  
 REFERENCE 1 (bases 1 to 11944)  
 AUTHORS Stephens, R.S., Kalman, S., Lammel, C.J., Fan, J., Marathe, R., Aravind, L., Mitchell, W.P., Olinger, L., Tatusov, R.L., Zhao, Q., Koonin, E.V. and Davis, R.W.  
 TITLE Genome sequence of an obligate intracellular pathogen of humans: Chlamydia trachomatis  
 JOURNAL Science 282 (5589), 754-759 (1998)  
 MEDLINE 99000809  
 PUBMED 97841136  
 REFERENCE 2 (bases 1 to 11944)  
 AUTHORS Kalman, S., Mitchell, W., Marathe, R., Lammel, C., Fan, J., Hyman, R.W., Olinger, L., Grimwood, J., Davis, R.W. and Stephens, R.S.  
 TITLE Comparative genomes of Chlamydia pneumoniae and C. trachomatis  
 JOURNAL Nat. Genet. 21 (4), 385-389 (1999)  
 MEDLINE 99206606  
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 REFERENCE 3 (bases 1 to 11944)  
 AUTHORS Stephens, R.S., Kalman, S., Lammel, C.J., Fan, J., Marathe, R., Aravind, L., Mitchell, W.P., Olinger, L., Tatusov, R.L., Zhao, Q., Koonin, E.V. and Davis, R.W.  
 TITLE Direct Submission  
 JOURNAL Submitted (20-MAY-1998) Program in Infectious Diseases, University of California, 235 Warren Hall, Berkeley, CA 94720-7360, USA  
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/db_xref="GI:3328602"
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5712..6728
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GETDKYARELGLPYPGOKLEELAREGDADAFSPARVSGVDFSFGLKTAVALY
KGNNSAKAPPEVSETOKRNIAASFQKAVPMTIAQKLDIVKTSFCSGLIVGGVAN
NSYFRLNLQICSLPIYFSSQLCSDNAMTAGLGRFLCNRTHSVSEKIVPCARYQNE
SACS"
5692..8248
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/sequence="CT198"
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/sequence="oppA_3"
/codon_start=1
/transl_table=11
/product="Oligopeptide Binding Protein"
/protein_id="AAC67790.1"
/db_xref="GI:3328604"

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TPHFLKILITLPVFPVHSQHOIRKEKESLPISTGAFFLKKKDRWLKLEKSPYYNK
DOVAOEICIHIPDOQTASALNOGKLDWGLPWGHSIQEETLATNNKRAPRSFDI
SQTSLFNTAKPFSHKLQALSLVNLKALASLAFVPAKHLPLPAHLHTYPEQPS
YKQENATLAKSLEALTELMTLEDEKYLPTFSATSTMSQIAQMLRDQWRESLG
ITFFPCREKVALLONDLIGNTFMSIGGFADFDPLAFISIFSSKGVKPYALODPQF
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LVMRYAKNS"
8407..9348
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8407..9348
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/codon_start=1
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/db_xref="GI:3328605"
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9380..10225
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9380..10225
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/db_xref="GI:3328606"
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RYSIALATAPPELLIADDESTALDSISQAVRLVITQIHQNSTALLTHNLAYS
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11066..11809

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Query Match 59.1% Score 26; DB 1; Length 11944;  
 Best Local Similarity 100.00; Pred. No. 4.2;  
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QY 19 ccatgcgcaagatcatcagtggaatc 44  
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 Db 6690 CCATGCGCAAGATATCATG7GGAATC 6715

RESULT 5  
 AX179663  
 LOCUS  
 DEFINITION Sequence 3 from Patent WO0146225.  
 AX179663 43 bp DNA linear PAT 06-AUG-2001

ACCESSION AX179663  
 VERSION AX179663.1 GI:15132084  
 KEYWORDS  
 SOURCE synthetic construct.  
 ORGANISM synthetic construct.  
 REFERENCE 1 (bases 1 to 43).  
 AUTHORS Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.  
 TITLE Chlamydia antigens and corresponding dna fragments and uses thereof  
 JOURNAL Patent: WO 0146225-A 3 28-JUN-2001;  
 Aventis Pasteur Limited (CA)  
 FEATURES Location/Qualifiers  
 source 1..43  
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 /note="5' PCR primer"  
 BASE COUNT 16 a 9 c 6 g 12 t  
 ORIGIN  
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 Best Local Similarity 81.1%; Pred. No. 2.7;  
 Matches 30; Conservative 0; Mismatches 7; Indels 0; Gaps 0;  
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 Db 1 ATAGATGCGCGCCGCCACCAATGAATAATTATTATT 37  
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 RESULT 6  
 AX278234  
 LOCUS AX278234 45 bp DNA linear PAT 01-NOV-2001  
 DEFINITION Sequence 3 from Patent WO0175114.  
 ACCESSION AX278234  
 VERSION AX278234.1 GI:16605283  
 KEYWORDS  
 SOURCE synthetic construct.  
 ORGANISM synthetic construct.  
 REFERENCE 1 (sites)  
 AUTHORS Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.  
 TITLE Chlamydia antigens and corresponding dna fragments and uses thereof  
 JOURNAL Patent: WO 0175114-A 3 11-OCT-2001;  
 Aventis Pasteur Limited (CA)  
 FEATURES Location/Qualifiers  
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 /note="5' PCR primer"  
 BASE COUNT 12 a 14 c 8 g 11 t  
 ORIGIN  
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 Best Local Similarity 93.1%; Pred. No. 2.7;  
 Matches 27; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 ataagaatgcggccgaccacatgcgaag 29  
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 Db 1 ATAGATGCGCGCCGCCACCAATGCAGAG 29  
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 RESULT 7  
 AX300411  
 LOCUS AX300411 45 bp DNA linear PAT 30-NOV-2001  
 DEFINITION Sequence 23 from Patent WO0185972.  
 ACCESSION AX300411  
 VERSION AX300411.1 GI:17381802  
 KEYWORDS  
 SOURCE synthetic construct.  
 ORGANISM synthetic construct.  
 REFERENCE 1 (sites)

AUTHORS Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.  
 TITLE Chlamydia antigens and corresponding dna fragments and uses thereof  
 JOURNAL Patent: WO 0185972-A 23 15-NOV-2001;  
 Aventis Pasteur Limited (CA)  
 FEATURES Location/Qualifiers  
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 /note="5' PCR primer"  
 BASE COUNT 13 a 11 c 13 g 8 t  
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 Best Local Similarity 100.0%; Pred. No. 5.9;  
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 ataagaatgcggccgaccacatgcg 25  
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 Db 1 ATAGATGCGCGCCGCCACCAATGCg 25  
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 RESULT 8  
 AX179705  
 LOCUS AX179705 46 bp DNA linear PAT 06-AUG-2001  
 DEFINITION Sequence 3 from Patent WO0146226.  
 ACCESSION AX179705  
 VERSION AX179705.1 GI:15132097  
 KEYWORDS  
 SOURCE synthetic construct.  
 ORGANISM synthetic construct.  
 REFERENCE 1 (bases 1 to 46)  
 AUTHORS Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.  
 TITLE Chlamydia antigens and corresponding dna fragments and uses thereof  
 JOURNAL Patent: WO 0146226-A 3 28-JUN-2001;  
 Aventis Pasteur Limited (CA)  
 FEATURES Location/Qualifiers  
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 /note="5' PCR primer"  
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 Matches 28; Conservative 0; Mismatches 5; Indels 0; Gaps 0;  
 QY 1 ataagaatgcggccgaccacatgcgaagatat 33  
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 Db 1 ATAGATGCGCGCCGCCACCAATGCACACATCT 33  
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 RESULT 9  
 AX146974  
 LOCUS AX146974 45 bp DNA linear PAT 08-JUN-2001  
 DEFINITION Sequence 3 from Patent WO0136455.  
 ACCESSION AX146974  
 VERSION AX146974.1 GI:14346245  
 KEYWORDS  
 SOURCE synthetic construct.  
 ORGANISM synthetic construct.  
 REFERENCE 1 (bases 1 to 45)  
 AUTHORS Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.  
 TITLE Chlamydia antigens and corresponding dna fragments and uses thereof  
 JOURNAL Patent: WO 0136455-A 3 25-MAY-2001;  
 Aventis Pasteur Limited (CA)  
 FEATURES Location/Qualifiers  
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ORIGIN

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Best Local Similarity 80.6%; Pred. No. 7.2;
Matches 29; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

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Db 1 ATAAGAATGCGCGCCACCATTGTAACAGTTTCAG 36
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RESULT 10
LOCUS      AX300423      45 bp      DNA      linear      PAT 30-NOV-2001
DEFINITION Sequence 35 from Patent WO0185972.
ACCESSION  AX300423
VERSION     AX300423.1 GI:17381814
KEYWORDS   Synthetic construct.
SOURCE      synthetic construct
ORGANISM    artificial sequence.
REFERENCE   1 (sites)
AUTHORS     Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.
TITLE       Chlamydia antigens and corresponding dna fragments and uses thereof
JOURNAL     Patent: WO 0185972-A 35 15-NOV-2001;
            Aventis Pasteur Limited (CA)
FEATURES   Location/Qualifiers
            source
            1..45
            /organism="synthetic construct"
            /db_xref="taxon:32630"
            /note="5' PCR primer"
BASE COUNT      13 a      14 c      10 g      8 t
ORIGIN

Query Match      56.4%; Score 24.8; DB 6; Length 45;
Best Local Similarity 92.9%; Pred. No. 7.2;
Matches 26; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ataagaatgcgcgcaccatgcgcaa 28
    |||||
Db 1 ATAAGAATGCGCGCCACCATTGCTA 28
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RESULT 11
LOCUS      AX300417      48 bp      DNA      linear      PAT 30-NOV-2001
DEFINITION Sequence 29 from Patent WO0185972.
ACCESSION  AX300417
VERSION     AX300417.1 GI:17381808
KEYWORDS   Synthetic construct.
SOURCE      synthetic construct
ORGANISM    artificial sequence.
REFERENCE   1 (sites)
AUTHORS     Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.
TITLE       Chlamydia antigens and corresponding dna fragments and uses thereof
JOURNAL     Patent: WO 0185972-A 29 15-NOV-2001;
            Aventis Pasteur Limited (CA)
FEATURES   Location/Qualifiers
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            1..48
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            /note="5' PCR primer"
BASE COUNT      17 a      11 c      8 g      12 t
ORIGIN

Query Match      56.4%; Score 24.8; DB 6; Length 48;
Best Local Similarity 92.9%; Pred. No. 7.2;
Matches 26; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ataagaatgcgcgcaccatgcgcaa 28
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Db 1 ATAAGAATGCGCGCCACCATTGCTA 28
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RESULT 12
LOCUS      AX147161      44 bp      DNA      linear      PAT 08-JUN-2001
DEFINITION Sequence 3 from Patent WO0136457.
ACCESSION  AX147161
VERSION     AX147161.1 GI:14346332
KEYWORDS   synthetic construct.
SOURCE      synthetic construct
ORGANISM    artificial sequence.
REFERENCE   1 (bases 1 to 44)
AUTHORS     Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.
TITLE       Chlamydia antigens and corresponding dna fragments and uses thereof
JOURNAL     Patent: WO 0136457-A 3 25-MAY-2001;
            Aventis Pasteur Limited (CA)
FEATURES   Location/Qualifiers
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            /db_xref="taxon:32630"
            /note="5' PCR primer"
BASE COUNT      15 a      17 c      7 g      5 t
ORIGIN

Query Match      55.5%; Score 24.4; DB 6; Length 44;
Best Local Similarity 96.2%; Pred. No. 11;
Matches 25; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ataagaatgcgcgcaccatgcgc 26
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Db 1 ATAAGAATGCGCGCCACCATTGCTA 26
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RESULT 13
LOCUS      AE002315      12173 bp      DNA      linear      BCT 26-MAY-2000
DEFINITION Chlamydia muridarum, section 46 of 85 of the complete genome.
ACCESSION  AE002315 AE002160
VERSION     AE002315.2 GI:8163226
KEYWORDS   Chlamydia muridarum.
SOURCE      Chlamydia muridarum
ORGANISM    Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
REFERENCE   1 (bases 1 to 12173)
AUTHORS     Read,T.D., Brunham,R., Shen,C., Gill,S.R., Heidelberg,J.F.,
            White,O., Hickey,E.K., Peterson,J., Umayam,L.A., Utterback,T.,
            Berry,K., Bass,S., Linher,K., Weidman,J., Khouri,H., Craven,B.,
            Bowman,C., Dodson,R., Gwinn,M., Nelson,W., DeBoy,R., Kolonay,J.,
            McClarty,G., Salzberg,S.L., Eisen,J. and Fraser,C.M.
            Genome sequences of Chlamydia trachomatis MoPu and Chlamydia
            pneumoniae AR39
            Nucleic Acids Res. 28 (6), 1397-1406 (2000)
TITLE       Chlamydia muridarum.
JOURNAL     Nucleic Acids Res. 28 (6), 1397-1406 (2000)
MEDLINE     20150255
PUBMED      10684935
REFERENCE   2 (bases 1 to 12173)
AUTHORS     Read,T.D., Brunham,R., Shen,C., Gill,S.R., Heidelberg,J.F.,
            White,O., Hickey,E.K., Peterson,J., Umayam,L.A., Utterback,T.,
            Berry,K., Bass,S., Linher,K., Weidman,J., Khouri,H., Craven,B.,
            Bowman,C., Dodson,R., Gwinn,M., Nelson,W., DeBoy,R., Kolonay,J.,
            McClarty,G., Salzberg,S.L., Eisen,J. and Fraser,C.M.
            Direct Submission
            Submitted (01-MAR-2000) The Institute for Genomic Research, 9712
            Medical Center Dr, Rockville, MD 20850, USA
            On Jun 1, 2000 this sequence version replaced gi:190506.
            Location/Qualifiers

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DELLPHRDPAFQSSQRTYAEWESKSLDYHLANFGYQSMYVHGHTFPDORKLGCQ
FVEDLPFGSAIGSKNLRIIVGVDTTANLSIERPHLIGLIDLSIPWATVGRF
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/db_xref="GI:8163227"
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NTAGRLMTNEFFALMETTVKEVATCIRNPPGDLTRLVFLVDFKGEIGQFTDRSLI
IASPEMLKQIMSPVSHKVLADTTREEVVDLVERKVALVPVVDENFLIGAITVEDV
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DEFINITION Sequence 3 from Patent WO0121803.
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SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 42)
AUTHORS Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.
TITLE I chlamydia /i antigens and corresponding dna fragments and uses
JOURNAL Patent: WO 0121803-A 3 29-MAR-2001;
Aventis Pasteur Limited (CA)
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RESULT 15
LOCUS AX268467 42 bp DNA linear PAT 29-OCT-2001
DEFINITION Sequence 3 from Patent WO0175113.
ACCESSION AX268467

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VERSION AX268467.1 GI:16541650
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (sites)
AUTHORS Murdin,A.D., Oomen,R.P., Wang,J. and Dunn,P.
TITLE Chlamydia antigens and corresponding dna fragments and uses thereof
JOURNAL Patent: WO 0175113-A 3 11-OCT-2001;
Aventis Pasteur Limited (CA)
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